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(19) (CA) APPLICATION FOR CANADIAN PATENT (12)

(54) Substituted Triazolinones

(72) Haas, Wilhelm - Germany (Federal Republic of); Findeisen, Kurt - Germany (Federal Republic of); Linker, Karl-Heinz - Germany (Federal Republic of); Schallner, Otto - Germany (Federal Republic of); König, Klaus - Germany (Federal Republic of); Marhold, Albrecht - Germany (Federal Republic of); Santel, Hans-Joachim - Germany (Federal Republic of); Dollinger, Markus - Germany (Federal Republic of); Wachendorff-Neumann, Ulrike - Germany (Federal Republic of);

- (71) Bayer Aktiengesellschaft Germany (Federal Republic of)
 ;
- (30) (DE) P 42 38 125.8 1992/11/12
- (57) 29 Claims

Notice: This application is as filed and may therefore contain an incomplete specification.

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Substitutes triazolinones

Abstract

The invention relates to new substituted triazolinones of the general formula (I)

in which

- R1 represents halogenoalkyl,
- R² represents hydrogen, amino, cyano, alkyl, alkenyl, alkinyl, halogenoalkyl, halogenoalkenyl, halogenoalkinyl, alkoxyalkyl, alkylideneimino, or in each case optionally substituted cycloalkyl or cycloalkylalkyl,
- R3 represents hydrogen or halogen,
- R4 represents cyano or nitro,
- R^5 represents nitro, cyano, halogen, heterocyclyloxy, a radical of the formula R^6 , $-O-R^6$, $-S-R^6$, $-S(0)-R^6$, $-SO_2-R^6$, $-SO_2-O-R^6$, $-O-SO_2-R^6$, $-C(0)-O-R^6$, $-NR^6R^7$, $-SO_2-NR^6R^7$, $-C(0)-NR^6R^7$, $-NH-P(0)(OR^6)(R^7)$ or $-NH-P(0)(OR^6)(R^7)$ or a radical of the formula

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and

- X represents oxygen or sulphur, where
- R⁶ and R⁷ independently of one another in each case represent hydrogen or in each case straight-chain or branched, optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl or aryl,

to a plurality of processes for their preparation, and to their use as herbicides, insecticides and acaricides.

The invention relates to new substituted triazolinones, to a plurality of processes for their preparation, and to their use as herbicides, insecticides and acaricides.

It has been disclosed that certain substituted triazolinones such as, for example, the compound 3,4-dimethyl-1-(3-fluoro-4-cyano-phenyl)-1,2,4-triazolin-5-one or the compound 3-methyl-4-propargyl-1-(2,5-difluoro-4-cyano-phenyl)-1,2,4-triazolin-5-one have herbicidal properties (cf., for example, DE 3,839,480).

However, the herbicidal activity of these previously known compounds against problem weeds as well as their compatibility with important crop plants are not entirely satisfactory in all fields of application.

New substituted triazolinones of the general formula (I)

15 in which

- R1 Pepresents halogenoalkyl,
- R² represents hydrogen, amino, cyano, alkyl, alkenyl, alkinyl, halogenoalkyl, halogeno-alkenyl, halogenoalkinyl, alkoxyalkyl, alkyl-ideneimino, or in each case optionally substituted cycloalkyl or cycloalkylalkyl,
- R3 represents hydrogen or halogen,
- R4 represents cyano or nitro,
- $R^5 \quad \text{represents nitro, cyano, halogen, hetero-} \\ 10 \quad \text{cyclylalkoxy, a radical of the formula } R^6, -O-R^6, \\ -S-R^6, -S(O)-R^6, -SO_2-R^6, -SO_2-O-R^6, -O-SO_2-R^6, \\ -C(O)-O-R^6, -NR^6R^7, -SO_2-NR^6R^7, -C(O)-NR^6R^7, \\ -NH-P(O)(OR^6)(R^7) \quad \text{or } -NH-P(O)(OR^6)(OR^7) \quad \text{or a radical of the formula}$



15 and

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- X represents oxygen or sulphur, where
- R⁶ and R⁷ independently of one another in each case represent hydrogen or in each case straight-chain or branched, optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl, cycloalkylalkyl, arylalkyl or aryl,

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have now been found.

Where appropriate, the compounds of the formula (I) can exist in the form of geometric and/or optical isomers or isomer mixtures of various compositions, depending on the nature of the substituents. The invention claims the pure isomers and the isomer mixtures.

Furthermore, it has been found that the new substituted triazolinones of the general formula (I)

in which

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- 10 R¹ represents halogenoalkyl,
 - R² represents hydrogen, amino, cyano, alkyl, alkenyl, alkinyl, halogenoalkyl, halogenoalkenyl, halogenoalkinyl, alkoxyalkyl, alkylideneimino, or in each case optionally substituted cycloalkyl or cycloalkylalkyl,
 - R3 represents hydrogen or halogen,

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R4 resents cyano or nitro,

R⁵ represents nitro, cyano, halogen, heterocyclylalkoxy, a radical of the formula R^6 , $-O-R^6$, $-S-R^6$, $-S(O)-R^6$, $-SO_2-R^6$, $-SO_2-O-R^6$, $-O-SO_2-R^6$, $-C(O)-O-R^6$, $-NR^5R^7$, $-SO_2-NR^6R^7$, $-C(O)-NR^6R^7$, $-NH-P(O)(OR^6)(R^7)$ or $-NH-P(O)(OR^6)(OR^7)$ or a radical of the formula

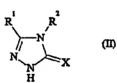
and

X represents oxygen or sulphur, where

10 R⁶ and R⁷ independently of one another in each case represent hydrogen or in each case straight-chain or branched, optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl, cycloalkylalkyl, arylalkyl or aryl,

are obtained when

15 a) 1H-triazolinones of the formula (II)



in which .

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R1, R2 and X have the abovementioned meanings,

are reacted with halogenobenzene derivatives of the formula (III)

5 in which

 R^3 , R^4 and R^5 have the abovementioned meanings and Hal represents halogen,

if appropriate in the presence of a diluent and if appropriate in the presence of a reaction auxiliary,

10 or when

b) substituted triazolinones of the formula (Ia)

in which

 R^1 , R^2 , R^3 , R^4 and X have the abovementioned meanings and

R5-1 represents halogen,

are reacted with nucleophiles of the formula (IV) ${\sf R^{6-1}\text{-}Z\text{-}H} \qquad \qquad ({\sf IV})$

in which

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Z represents oxygen or sulphur and

R⁶⁻¹ represents in each case straight-chain or branched, optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl or aryl, and furthermore, in the event that Z represents oxygen, R⁶⁻¹ also represents heterocyclyl,

if appropriate in the presence of a diluent and if appropriate in the presence of a reaction auxiliary, or when

c) substituted triazolinones of the formula (Ib)

in which

 R^1 , R^3 , R^4 , R^5 and X have the abovementioned meanings and

R2-1 represents amino,

are reacted with sodium nitrite in the presence of an acid and, if appropriate, in the presence of a diluent, or when

d) substituted triazolinones of the formula (Ic)

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in which



 R^1 , R^3 , R^4 , R^5 and X have the abovementioned meanings and

R2-2 represents hydrogen,

are reacted with alkylating agents of the formula (V)

 $R^{2-3}-E \qquad \qquad (V)$

in which

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R²⁻³ represents alkyl, alkenyl, alkinyl, halogenoalkyl, halogenoalkenyl, halogenoalkinyl, alkoxyalkyl or optionally substituted cycloalkyl and

E represents an electron-attracting leaving group,

if appropriate in the presence of a diluent and if appropriate in the presence of a reaction auxiliary.

Finally, it has been found that the new substituted triazolinones of the general formula (I) have herbicidal, insecticidal and acaricidal properties.

Surprisingly, the substituted triazolinones of the general formula (I) according to the invention have a considerably better herbicidal activity against problem weeds and unexpectedly, at the same time, also a considerably better acaricidal activity compared with the

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substituted triazolinones known from the prior art such as, for example, the compound 3,4-dimethyl-1-(3-fluoro-4-cyano-phenyl)-1,2,4-triazolin-5-one or the compound 3-methyl-4-propargyl-1-(2,5-difluoro-4-cyano-phenyl)-1,2,4,triazolin-5-one, which are similar compounds-chemically and from the point of view of their action.

Formula (I) provides a general definition of the substituted triazolinones according to the invention. Preferred compounds of the formula (I) are those in which

- R¹ represents straight-chain or branched halogenoalkyl having 1 to 6 carbon atoms and 1 to 13 identical or different halogen atoms, in particular fluorine, chlorine, bromine or iodine,
- represents hydrogen, amino, cyano, straight-chain or \mathbb{R}^2 branched alkyl having 1 to 8 carbon atoms, in each 15 case straight-chain or branched alkenyl or alkinyl, each of which has 2 to 6 carbon atoms, straightchain or branched halogenoalkyl having 1 to 6 carbon atoms and 1 to 13 identical or different halogen atoms, in particular fluorine, chlorine, bromine or 20 iodine, in each case straight-chain or branched halogenoalkenyl or halogenoalkinyl, each of which has 2 to 6 carbon atoms and 1 to 11 identical or different halogen atoms, in particular fluorine, chlorine, bromine or iodine, straight-chain or 25 branched alkoxyalkyl having 1 to 4 carbon atoms in each of the individual alkyl moieties, straight-chain

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or branched alkylideneimino having 1 to 8 carbon atoms, or cycloalkyl or cycloalkylalkyl, each of which has 3 to 8 carbon atoms in the cycloalkyl moiety and, if appropriate, 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, and each of which is optionally monosubstituted or polysubstituted in the cycloalkyl moiety by identical or different halogen substituents, in particular fluorine, chlorine, bromine and/or iodine,

- 10 R³ represents hydrogen, fluorine, chlorine, bromine or iodine,
 - R4 represents cyano or nitro,
- R⁵ represents nitro, cyano, fluorine, chlorine, bromine, iodine or heterocyclyl $-C_1-C_4$ -alkoxy, the hetero-15 cyclyl radical being represented by a three- to sevenmembered, optionally benzo-fused, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms, in particular nitrogen, oxygen and/or sulphur, or a radical of the formula 20 R^{6} , $-O-R^{6}$, $-S-R^{6}$, $-S(O)-R^{6}$, $-SO_{2}-R^{6}$, $-SO_{2}-O-R^{6}$, $-O-SO_2-R^6$, $-C(O)-O-R^6$, $-NR^6R^7$, $-SO_2-NR^6R^7$, $-C(O)-NR^6R^7$, $-NH-P(O)(OR^6)(R^7)$ or $-NH-P(O)(OR^6)(OR^7)$ or a radical of the formula

25 and

X represents oxygen or sulphur, where

R⁶ and R⁷ independently of one another in each case represent hydrogen or straight-chain or branched alkyl which has 1 to 8 carbon atoms and which is optionally monosubstituted or polysubstituted by identical or different substituents, suitable substituents being:

halogen, in particular fluorine, chlorine, bromine and/or iodine, cyano, carboxyl, carbamoyl, in each case straight-chain or branched alkoxy, alkoxy-alkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, alkoxycarbonylalkyl, N-alkylamino-carbonyl, cycloalkylaminocarbonyl, N,N-dialkylaminocarbonyl, trialkylsilyl or alkylsulphonylaminocarbonyl, each of which has 1 to 8 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl being represented by a five- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms, in particular nitrogen, oxygen and/or sulphur;

R⁶ and R⁷ furthermore represent alkenyl or alkinyl, each of which has 2 to 8 carbon atoms and each of which is optionally monosubstituted or polysubstituted by identical or different halogen substituents, in particular fluorine, chlorine, bromine and/or iodine;

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R⁶ and R⁷ furthermore represent cycloalkyl which has 3 to 7 carbon atoms and which is optionally monosubstituted or polysubstituted by identical or different halogen substituents, in particular fluorine, chlorine, bromine and/or iodine, and/or by straight-chain or branched alkyl having 1 to 4 carbon atoms, or represent C₂-C₇-cycloalkyl-C₁-C₃-alkyl, or

R⁶ and R⁷ represent arylalkyl or aryl, each of which has 6 to 10 carbon atoms in the aryl moiety and, if appropriate, 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, and each of which is optionally monosubstituted or polysubstituted in the aryl moiety by identical or different substituents, suitable aryl substituents in each case being:

halogen, cyano, nitro, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl, each of which has 1 to 6 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl, each of which has 1 to 6 carbon atoms and 1 to 13 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl, each of which has 1 to 6 carbon atoms in the individual alkyl moieties, and phenyl which is optionally monosubstituted or polysubstituted by identical or different halogen substituents and/or by straight-chain or branched alkyl

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or alkoxy, each of which has 1 to 6 carbon atoms, and/or by straight-chain or branched halogenoalkyl or halogenoalkoxy, each of which has 1 to 6 carbon atoms and 1 to 13 identical or different halogen atoms.

Particularly preferred compounds of the formula (I) are those in which

- R¹ represents straight-chain or branched halogenoalkyl having 1 to 4 carbon atoms and 1 to 9 identical or different halogen atoms, in particular fluorine, chlorine or bromine.
- R2 represents hydrogen, amino, cyano, straight-chain or branched alkyl having 1 to 6 carbon atoms, in each case straight-chain or branched alkenyl or alkinyl, 15 each of which has 2 to 4 carbon atoms, straightchain or branched halogenoalkyl having 1 to 4 carbon atoms and 1 to 9 identical or different halogen atoms, in particular fluorine, chlorine or bromine, in each case straight-chain or branched halogeno-20 alkenyl or halogenoalkinyl, each of which has 2 to 4 carbon atoms and 1 to 7 identical or different halogen atoms, in particular fluorine, chlorine or bromine, straight-chain or branched alkoxyalkyl having 1 to 3 carbon atoms in each of the individual 25 alkyl moieties, straight-chain or branched alkylideneimino having 1 to 6 carbon atoms, or cycloalkyl or cycloalkylalkyl, each of which has 3 to 7 carbon

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atoms in the cycloalkyl moiety and, if appropriate, 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety, and each of which is optionally monosubstituted to tetrasubstituted in the cycloalkyl moiety by identical or different halogen substituents, in particular fluorine, chlorine and/or bromine,

- R³ represents hydrogen, fluorine, chlorine or bromine,
- R4 represents cyano or nitro,
- 10 R⁵ represents nitro, cyano, fluorine, chlorine, bromine or heterocyclyl -C₁-C₃-alkoxy, the heterocyclyl radical being represented by a four- or six-membered, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms, in particular nitrogen, oxygen and/or sulphur, or a radical of the formula R⁶, -O-R⁶, -S-R⁶, -S(O)-R⁶, -SO₂-R⁶, -SO₂-O-R⁶, -O-SO₂-R⁶, -C(O)-O-R⁶, -NR⁶R⁷, -SO₂-NR⁶R⁷, -C(O)-NR⁶R⁷, -NH-P(O)(OR⁶)(R⁷) or -NH-P(O)(OR⁶)(OR⁷) or a radical of the formula

R¹ R²

and

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X represents oxygen or sulphur, where

R6 and R7 independently of one another in each case

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represent hydrogen or straight-chain or branched alkyl which has 1 to 6 carbon atoms and which is optionally monosubstituted, suitable substituents being:

cyano, carboxyl, carbamoyl, in each case straightchain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, alkoxycarbonylalkyl, N-alkylaminocarbonyl, dialkylaminocarbonyl, trialkylsilyl alkylsulphonylaminocarbonyl, each of which has 1 to 6 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl radical being represented by a five- or six-membered, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms, in particular nitrogen, oxygen and/or sulphur;

R⁶ and R⁷ furthermore represent straight-chain or branched halogenoalkyl having 1 to 4 carbon atoms and 1 to 9 identical or different halogen atoms, in particular fluorine, chlorine or bromine and being optionally further substituted by C₁₋₂alkoxycarbonyl, C₁₋₆cycloalkylaminocarbonyl or cyano, R⁶ and R⁷ furthermore represent alkenyl or alkinyl, each of which has 2 to 6 carbon atoms and each of which is optionally monosubstituted to trisubstituted by identical or different halogen substituents, in particular fluorine, chlorine or bromine;

R6 and R7 furthermore represent cycloalkyl which has 3 to

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6 carbon atoms and which is optionally monosubstituted to tetrasubstituted by identical or different halogen substituents, in particular fluorine, chlorine or bromine, and/or by straight-chain or branched alkyl having 1 to 3 carbon atoms, or represent C_{3-6} -cycloalkyl- C_1 - C_2 -alkyl, or

represent phenylalkyl or phenyl, the first-mentioned has 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety and each of which is optionally monosubstituted to trisubstituted in the phenyl moiety by identical or different substituents, suitable phenyl substituents in each case being:

halogen, cyano, nitro, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl, each of which has 1 to 4 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl, each of which has 1 to 4 carbon atoms and 1 to 9 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl, each of which has 1 to 4 carbon atoms in the individual alkyl moieties, and phenyl which is optionally monosubstituted or polysubstituted by identical or different halogen substituents and/or by straight-chain or branched alkyl or alkoxy, each of which has 1 to 4 carbon atoms, and/or straight-chain or branched halogenoalkyl

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halogenoalkoxy, each of which has 1 to 4 carbon atoms and 1 to 9 identical or different halogen atoms.

Very particularly preferred compounds of the formula (I) are those in which

- R¹ represents halogenoalkyl having 1 or 2 carbon atoms and 1 to 5 identical or different halogen atoms, in particular fluorine or chlorine.
- \mathbb{R}^2 represents hydrogen, amino, cyano, straight-chain or 10 branched alkyl having 1 to 4 carbon atoms, in each case straight-chain or branched alkenyl or alkinyl, each of which has 2 to 3 carbon atoms, halogenoalkyl having 1 or 2 carbon atoms and 1 to 5 identical or different halogen atoms, in particular fluorine, 15 chlorine or bromine, in each case straight-chain or branched halogenoalkenyl or halogenoalkinyl, each of which has 2 to 3 carbon atoms and 1 to 3 identical or different halogen atoms, in particular fluorine or chlorine, straight-chain or branched alkoxyalkyl 20 having 1 or 2 carbon atoms in each of the individual alkyl moieties, straight-chain or branched alkylideneimino having 1 to 6 carbon atoms, or cyclopropyl, cyclopropylmethyl, cyclohexyl or cyclohexylmethyl, each of which is optionally monosubstituted 25 or disubstituted in the cycloalkyl moiety by identical or different halogen substituents, in particular fluorine or chlorine,



- R3 represents hydrogen, fluorine or chlorine,
- R4 represents cyano or nitro,
- represents nitro, cyano, fluorine, chlorine, bromine or heterocyclylmethoxy, the heterocyclyl radical being represented by a five- or six-membered, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms, in particular nitrogen, oxygen and/or sulphur, or represents a radical of the formula R⁶, -O-R⁵, -S-R⁶, -S(0)-R⁶, -SO₂-R⁶, -SO₂-R⁶, -C(0)-O-R⁶, -NR⁶R⁷, -SO₂-NR⁶R⁷, -C(0)-NR⁶R⁷, -NH-P(0)(OR⁶)(R⁷) or -NH-P(0)(OR⁶)(OR⁷) or a radical of the formula



and

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- 15 X represents oxygen or sulphur, where
 - R⁵ and R⁷ independently of one another in each case represent hydrogen or optionally monosubstituted straight-chain or branched alkyl having 1 to 4 carbon atoms, suitable substituents being:
- cyano, carboxyl, carbamoyl, in each case straightchain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, alkylcarbonylalkyl, N-alkylaminocarbonyl,

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N.N-dialkylaminocarbonyl, trialkylsilyl or alkylsulphonylaminocarbonyl, each of which has 1 to
4 carbon atoms in the individual alkyl moieties, or
heterocyclyl, the heterocyclyl radical being represented by a five- or six-membered saturated or
unsaturated heterocycle having 1 to 3 identical or
different hetero atoms, in particular nitrogen,
oxygen and/or sulphur;

R⁶ and R⁷ furthermore represent halogenoalkyl having 1 or 2 carbon atoms and 1 to 5 identical or different halogen atoms, in particular fluorine or chlorine and being optionally further substituted by methoxycarbonyl, ethoxycarbonyl, cyano or cyclopacylaminocarbonyl;

R⁶ and R⁷ furthermore represent alkenyl or alkinyl, each of which has 2 to 5 carbon atoms and each of which is optionally monosubstituted by halogen, in particular fluorine or chlorine;

R⁵ and R⁷ furthermore represent cyclopropyl or cyclohexyl, each of which is optionally monosubstituted or disubstituted by identical or different substituents from the series comprising fluorine, chlorine, methyl and/or ethyl, or represent cyclopropylmethyl, cyclopentylmethyl or cyclohexylmethyl, or

R⁵ and R⁷ represent phenylalkyl or phenyl, the first-mentioned has 1 or 2 carbon atoms in the alkyl moiety and each of which is optionally monosubstituted or disubstituted in the phenyl moiety by identical or different substituents, suitable phenyl substituents in each case being:

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, chlorine, bromine, cyano, ni ethyl, n- or i-propyl, n-, i-, s- or t-butyl, methoxy, ethoxy, n- or i-propoxy, n-, i-, s- or tbutoxy, methylthio, ethylthio, methylsulphinyl, methylsulphonyl, trifluoromethyl, difluoromethyl, trifluoromethoxy, difluoromethoxy, trifluoromethyltrfluoromethylsulphinyl, trifluoromethylthio, sulphonyl, methoxycarbonyl, ethoxycarbonyl, methoximinomethyl, methoximinoethyl, ethoximinomethyl, ethoximinoethyl, or phenyl which is optionally monosubstituted to disubstituted by identical or different substituents from the series comprising fluorine, chlorine, bromine, methyl, ethyl, methoxy, ethoxy, trifluoromethyl and/or trifluoromethoxy.

The following substituted triazolinones of the general formula (I) may be mentioned individually in addition to the compounds given in the Preparation Examples:

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	R¹	R²	R³	R ⁴	R ⁵	x
	CF ₃	СН3	F	CN	ОН	0
	CF ₃	CH ₃	а	CN	ОН .	0
	CF ₃	CH ₃	F	NO_2	ОН	0
	CF ₃	CH ₃	а	NO ₂	ОН	0
	CF ₃	CH ₃	a	CN	CH ₃ O	0
	CF ₃	СН3	a	NO ₂	CH ₃ O	0
	CF ₃	CH ₃	F	NO ₂	CH ₃ O	0
	CF ₃	CH ₃	F	CN	-O-CH ₂ -C≡CH	O
	CF ₃	CH ₃	а	CN	-O-CH ₂ -C≡CH	0
5	CF ₃	CH ₃	F	NO ₂	-O-CH ₂ -C≡CH	0
•	CF ₃	CH ₃	a	NO ₂	-O-CH ₂ -C≡CH	0
	CF ₃	CH ₃	F	CN	-0CH-COOC ⁵ H ²	0
	CF ₃	CH ₃	a	CN	-0CH-COOC ² H²	0
	CF ₃	CH3	F	NO ₂	осн-соос ^у н Сн	0
	CF ₃	CH ₃	a	NO ₂	CH, −0−CH−COOC₂H,	Ο,

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R1	R²	R³	R ⁴	R ⁵	x
CF ₃	СН3	F	CN	 0.	0
CF ₃	СН3	CI	CN		O
CF ₃	СН3	F	NO ₂	-0-	0
CF3	CH ₃	а	NO ₂	-0-00	0
CF ₃	СН3	F	CN	-0-CH ₂ -CN	0
CF ₃	СН3	CI	CN	-O-CH ₂ -CN	0
CF ₃	СН3	F	NO ₂	-O-CH ₂ -CN	0
CF ₃	СН3	a	NO ₂	-O-CH ₂ -CN	0
CF ₃	CH ₃	F	CN	-0-CH2 0	0
CF ₃	CH ₃	a	CN	-0-CH ₂	0

R1	R²	R³	R ⁴	R ⁵	X
CF ₃	CH ₃	F	NO ₂	-0-CH2 0.	0
CF3	СН3	а	NO ₂	-0-CH ₂ 0	0
CF3	CH ₃	F	CN	CI O-CH ₂ -CH=CH ₄	0
CF ₃	СН3	а	CN	-0-CH-CH-CH' CI	0
CF ₃	СН3	а	NO ₂	o-ch-ch-ch c	0
CF3	СН3	F	NO ₂	0-CH²-CH=CH² CI	0
CF ₃	СН3	ŗ	CN	-O-SO ₂ -CH ₃	0
CF ₃	СН3	CI	CN	-O-SO ₂ -CH ₃	0
CF ₃	CH ₃	F	NO_2	-O-SO ₂ -CH ₃	O
CF ₃	СН3	a	NO ₂	-O-SO ₂ -CH ₃	O
CF ₃	СН3	F	CN	OCH ₂ -COOCH,	0

R1	R²	R³	R ⁴	R ⁵	x
CF ₃	СН3	a	CN	-0-СН,-СООСН,	0
CF ₃	СН3	F .	NO ₂	-0-CH-COOCH	0
CF3	CH ₃	a	NO ₂	-0-CH ₂ -COOCH ₃	0
CF ₃	СН3	F	NO ₂	F	0
CF ₃	СН3	a	NO ₂	F	0
CF3	СН3	F	NO ₂	a	0
CF ₃	CH ₃	а	NO ₂	а	0
CF ₃	CH ₃	F	CN	-O-CHF2	o
CF ₃	СН3	a	CN	-O-CHF ₂	0
CF ₃	CH ₃	F	NO_2	-O-CHF ₂	o
CF ₃	CH ₃	а	NO ₂	-0-CHF2	0
CF ₃	СН3	F	CN	-S-CH ₃	0
CF ₃	CH ₃	a	CN	-S-CH ₃	0
CF3	CH ₃	F	NO ₂	-S-CH ₃	0
CF ₃	CH ₃	а	NO_2	-S-CH ₃	0



R¹	R²	R³	R ⁴	R ⁵	x
CF ₃	CH ₃	F	CN	-S-C ₂ H ₅	o
CF ₃	CH ₃	a .	CN	-S-C ₂ H ₅ .	o
CF ₃	СН3	F	NO ₂	-S-C ₂ H ₅	o
CF3	CH ₃	а	NO ₂	-S-C ₂ H ₅	0
CF ₃	CH ₃	F	CN	СН, -s-сн-соос _г н,	0
CF ₃	СН3	а	CN	СН, —s—сн—соос,н,	0
CF ₃	CH ₃	F	NO ₂	СН, SСНСООС,Н,	0
CF ₃	СН3	а	NO ₂	СН ₃ —S—СН—СООС ₂ Н ₃	0
CF3	CH ₃	F	CN	CH _s −s−CH−C≡CH	0
CF ₃	CH ₃	α	CN	-S-CH-C≡CH	0
CF ₃	СН3	F	NO ₂	-2— CH -C≡CH	0
CF ₃	СН3	CI	NO ₂	CH, -S-CH-C≡CH	0

210	5 Ø
75	v

R1	R²	R ³	R ⁴	R ⁵	x
CF3	СН3	a	CN	-S-CH ₂ -COOCH ₃	o
CF ₃	CH ₃	F	CN	-S-CH ₂ -COOCH ₃	0
CF ₃	СН3	а	NO ₂	-S-CH ₂ -COOCH ₃	0
CF ₃	CH ₃	F	NO ₂	-S-CH ₂ -COOCH ₃	0
CF3	СН3	F	CN	CH ₃	o
CF ₃	СН3	F	CN	-S(O)-CH ₃	0
CF3	СН3	F	CN	-SO ₂ -CH ₃	0
CF ₃	CH ₃	F	CN	-SO ₂ -O-CH ₃	0
CF ₃	CH ₃	F	CN	-SO ₂ -NH-CH ₃	0
CF ₃	CH ₃	F	CN	-NH-CH ₃	0
CF ₃	CH ₃	F ·	CN	-N(CH ₃) ₂	0
CF ₃	CH ₃	F	CN	-COOCH3	0
CF ₃	СН3	F	CN	-COOC ₂ H ₅	0
CF ₃	CH ₃	a	NO ₂	-соосн ₃	o
CF ₃	CH ₃	a	NO ₂	-COOC ₂ H ₅	0
CF ₃	СН3	F	CN	-CO-NH-CH3	0
CF ₃	СН3	F	CN	-CO-N(CH ₂)-CH ₂	0

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R ¹	R ²	R³	R ⁴	R ⁵	x
CF3	CH ₃	F	CN	O NH-P OCH, CH,	0
CF ₃	СН3	F	CN	O NH-POC ₂ H ₃ OC ₂ H ₃	o
CF ₃	C_2H_5	F	CN	ОН	0
CF ₃	C ₂ H ₅	F	CN	OCH ₃	0
CF ₃	C ₂ H ₅	F	CN	-O-CH ₂ -CH=CH ₂	0
CF ₃	C ₂ H ₅	F	CN	-O-CH ₂ -C≡CH	0
CF ₃	C ₂ H ₅	F	CN	-о-сн-соос ⁵ н²	0
CF ₃	C ₂ H ₅	F	CN	-0-CH ₂ -COOCH ₃	0
CF ₃	C ₂ H ₅	F	CN	-S-CH ₃	O
CF ₃	C ₂ H ₅	F	CN	-S-C ₂ H ₅	·O .
CF ₃	C ₂ H ₅	F	CN	CH, -SCHCOOCH,	0
CF ₃	C ₂ H ₅	F	CN	F	0

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R ¹	R ²	R³	R ⁴	R ⁵	x
CF ₃	C ₂ H ₅	F	CN	-0-CH, 0.	0
CF ₃	C ₂ H ₅	F	CN	-0 CE-CE-CH -0 CE-CH	0
CF ₃	C ₂ H ₅	а	CN	OCH ₃	0
CF ₃	C ₂ H ₅	а	CN	-S-C ₂ H ₅	Ο
CF ₃	C ₂ H ₅	F	NO ₂	осн3	0
CF ₃	C ₂ H ₅	a	NO ₂	-O-CH ₂ -C≡CH	Ο
CF ₃	C ₂ H ₅	а	CN	oCHCOOCH, CH3	0
CF3	C ₂ H ₅	F	NO ₂	-о-сн-соосн _э	0
CF ₃	C ₂ H ₅	a a	NO ₂	-0-CH-COOCH ²	0
CF ₃	C ₂ H ₅	F	CN		0
CF ₃	C ₂ H ₅	F	CN	-O-СН ₂ -С _б Н ₅	0
CF ₃	-CH ₂ -CH=CH ₂	F	CN	ОН	0
CF ₃	-CH ₂ -CH=CH ₂	F	CN	OCH ₃	ο.

	R1	R ²	R³	R ⁴	R ⁵	x
	CF ₃	-CH ₂ -CH=CH ₂	F	CN	-O-CH ₂ -CH=CH ₂	O
	CF ₃	-CH ₂ -CH=CH ₂	F	CN	-O-CH ₂ -C≡CH·	0
	CF ₃	-CH ₂ -CH=CH ₂	F	CN	CH ₃ -O-CH-COOC ₂ H ₅	0
	CF ₃	-CH ₂ -CH=CH ₂	F	CN	O-CH ₂ -COOCH ₃	0
	CF ₃	-CH ₂ -CH=CH ₂	F	CN	-S-CH ₃	0
	CF3	-СH ₂ -СН=СН ₂	F	CN	-S-C ₂ H ₅	0
	CF ₃	-CH ₂ -CH=CH ₂	F	CN	CH, -SCH-COOCH,	0
	CF ₃	-CH ₂ -CH=CH ₂	F	CN	F	0
5	CF ₃	-CH ₂ -CH=CH ₂	F	CN	-0-CH ₂	0
3	CF ₃	-CH ₂ -CH=CH ₂	F	CN	-०-वम-ट≡वस वस	0
	CF ₃	-CH ₂ -CH=CH ₂	a	CN	OCH ₃	0
	CF ₃	-CH ₂ -CH=CH ₂	a	CN	-S-C ₂ H ₅	0
	CF ₃	-CH ₂ -CH=CH ₂	F	NO ₂	OCH ₃	0
	CF ₃	-CH ₂ -CH=CH ₂	a	NO ₂	-O-CH ₂ -C≡CH	0

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R1	R²	R³	R ⁴	R ⁵	x
CF ₃	-CH ₂ -CH=CH ₂	а	CN	-0-CH-COOCH,	0
CF ₃	-CH ₂ -CH=CH ₂	F	NO ₂	-0-CH-000CH,	0
CF ₃	-CH ₂ -CH=CH ₂	Cl	NO ₂	-о-сн-соосн, сн,	0
CF ₃	-CH ₂ -CH=CH ₂	F	CN	-0	0
CF ₃	-CH ₂ -CH=CH ₂	F	CN	-O-СH ₂ -С ₆ H ₅	0
CF ₃	-CHF ₂	F	CN	ОН	0
CF ₃	-CHF ₂	F	CN	OCH ₃	o
CF ₃	-CHF ₂	F	CN	-O-CH ₂ -CH=CH ₂	o
CF ₃	-CHF ₂	F	CN	-0-СН2-С≡СН	0
CF ₃	-CHF ₂	F	CN	-о-сн-соос'н² сн³	0
·CF ₃	-CHF ₂	F	CN	-0-CH ₂ -COOCH ₃	0
CF ₃	-CHF ₂	F	CN	-S-CH ₃	0
CF ₃	-CHF ₂	F	CN	-S-C ₂ H ₅	0

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4. .

R1	R ²	R³	R ⁴	R ⁵	x
CF ₃	-CHF ₂	F	CN	-S-CH-COOCH,	0
CF ₃	-CHF ₂	F	CN	F	0
CF3	-CHF ₂	F	CN	-0-CH ₂	0
CF3	-CHF ₂	F	CN	-0-CH-C≡CH CH	0
CF ₃	-CHF ₂	а	CN	OCH ₃	0
CF ₃	-CHF ₂	CI	CN	-S-C ₂ H ₅	0
CF ₃	-CHF ₂	F	NO ₂	OCH ₃	0
CF ₃	-CHF ₂	a	NO ₂	-О-СН ₂ -С≡СН	0
CF ₃	-CHF ₂	а	CN	-0-CH-COOCH,	0
CF ₃	-CHF ₂	F	NO ₂	-о-сн-соосн, сн,	0
CF ₃	-CHF ₂	а	NO ₂	-о-сн-соосн, сн,	0
CF ₃	-CHF ₂	F	CN		0

	R¹	R²	R³	R ⁴	R ⁵	x
	CF ₃	-CHF ₂	F	CN	-O-CH ₂ -C ₆ H ₅	ο
	CF ₃	$\overline{}$	F	CN	ОН .	0
	CF ₃	$\overline{}$	F	CN	OCH ₃	0
	CF ₃	$\overline{}$	F	CN	-O-CH ₂ -CH=CH ₂	0
	CF ₃	$\overline{}$	F	CN	-O-CH ₂ -C≡CH	0
	CF ₃	$\overline{}$	F	CN	ocHcooc³H² CH²	0
	CF ₃	$\overline{}$	F	CN	-0СН ₂ -соосн ₄	0
5	CF ₃	\longrightarrow	F	CN	-S-CH ₃	0
	CF ₃	$\overline{}$	F	CN	-S-C ₂ H ₅	0
	CF3		F	CN	_s_ ch, _s_ ch_ coch,	0
	CF ₃	$\overline{}$	F	CN	F	0

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R¹	R ²	R³	R ⁴	R ⁵	x
CF3	$\overline{}$	F	CN .	-0-CH ₂ 0.	0
CF ₃	$\overline{}$	F	CN	–0 − αਜ−ਟ≡਼ੁੁੁੁੁੁੁੁਰਮ ਰਸੰ	0
CF ₃	$\overline{}$	a	CN	OCH ₃	0
CF3	$\overline{}$	a	CN	-S-C ₂ H ₅	0
CF ₃	$\overline{}$	F	NO ₂	осн ₃	0
CF ₃	$\overline{}$	a	NO ₂	-O-CH ₂ -C≡CH	O
CF ₃	$\overline{}$	а	CN	СН, -0-СН-СООСН,	0
CF ₃	$\overline{}$	F	NO ₂	-0- СН -СООСН,	0
CF3	$\overline{}$	CI	NO ₂	-0CHCOOCH CH,	0
CF ₃	$\overline{}$	F	CN		o

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	R¹	R ²	R ³	R ⁴	R ⁵	x
	CF ₃	$\overline{}$	F	CN	-0-СН ₂ -С _б Н ₅	0
	-CHF ₂	CH ₃	F	CN	ОН	0
	-CHF ₂	СН3	F	CN	OCH ₃	0
	-CHF ₂	СН3	F	CN	-O-CH ₂ -CH=CH ₂	0
	-CHF ₂	СН3	F	CN	-О-СН2-С≡СН	0
	-CHF ₂	СН3	F	CN	_O_CH_COOC,H,	0
	-CHF ₂	СН3	F	CN	_о-сн ₂ -соосн ₃	0
	-CHF ₂	СН3	F	CN	-S-CH ₃	0
	-CHF ₂	CH ₃	F	CN	-S-C ₂ H ₅	0
5	-CHF ₂	CH ₃	F	CN	CH ₃ - -SCH-COOCH ₃	0
	-CHF ₂	СН3	F	CN	F	0
	-CHF ₂	СН3	F	CN	-0-CH ₂	0
	-CHF ₂	CH ₃	F	CN	CH, -0-CH-C≡CH	0

R1	R ²	R³	R ⁴	R ⁵	x
-CHF ₂	CH ₃	a	CN	OCH ₃	О
-CHF ₂	СН3	a	CN	-S-C ₂ H ₅	. 0
-CHF ₂	CH ₃	F	NO ₂	OCH ₃	0
-CHF ₂	CH ₃	CI	NO ₂	-O-CH ₂ -C≡CH	O
-CHF ₂	CH ₃	а	CN	-0-CH-COOCH,	0
-CHF ₂	CH ₃	F	NO ₂	-0-CH-COOCH,	0
-CHF ₂	СН3	а	NO ₂	-0-сн-соосн,	0
-CHF ₂	СН3	F	CN		0
-CHF2	СН3	F	CN	-O-CH ₂ -C ₆ H ₅	0
-CF ₂ CI	CH ₃	F	CN	OH	О
-CF ₂ CI	CH ₃	F	CN	OCH ₃	0
-CF ₂ CI	CH ₃	F	CN	-O-CH ₂ -CH=CH ₂	0
-CF ₂ CI	CH ₃	F	CN	-O-CH ₂ -C≡CH	O
-CF ₂ Cl	СН3	F	CN	СН ₃ ОСНСООС,Н ₄	0

R ¹	R²	R ³	R ⁴	R ⁵	x
-CF ₂ Cl	СН3	F	CN	0СН,-СООСН,	0
-CF ₂ CI	СН3	F	CN	-S-CH ₃	0
-CF ₂ CI	CH ₃	F	CN	-S-C ₂ H ₅	ο
-CF ₂ CI	CH3	F	CN	сн, -s-сн-соосн,	0
-CF ₂ CI	СН3	F	CN	F	0
-CF ₂ CI	СН3	F	CN	-0-CH ₂	0
-CF ₂ CI	СН3	F	CN	-o-a:-c≡an an²	0
-CF ₂ CI	CH ₃	a .	CN	OCH ₃	0
-CF ₂ CI	СН3	а	CN	-S-C ₂ H ₅	0
-CF ₂ CI	СН3	F	NO ₂	OCH ₃	0
-CF ₂ CI	CH ₃	a	NO ₂	-O-CH ₂ -C≡CH	0
-CF2CI	СН3	а	CN	CH, OCHCOOCH,	0



	R1	R²	R³	R ⁴	R ⁵	x
	-CF ₂ CI	CH ₃	F	NO ₂	-0-СН-СООСН ₃	0
	-CF ₂ CI	CH ₃	а	NO ₂	СН, ОСНСООСН,	.0
	-CF ₂ CI	СН3	F	CN	_0	0
	-CF ₂ CI	СН3	F	CN	-O-CH ₂ -C ₆ H ₅	0
	-CC1 ₃	CH ₃	F	CN	ОН	O
	-CCI ₃	CH ₃	F	CN	осн ₃	Ο
	-CCl ₃	СН3	F	- CN	-O-CH ₂ -CH=CH ₂	0
	-CCI ₃	СН3	F	CN	-O-CH ₂ -C≡CH	0
	-CC13	CH ₃	F	CN	-о-сн-соос ^т н	0
	-CCI ₃	CH ₃	F	CN	-0-CH ₂ -COOCH ₃	0
5	-CCI3	CH ₃	F	CN	-S-CH ₃	0
	-CCl ₃	CH ₃	F	CN	-S-C ₂ H ₅	0
	-CCl ₃	CH ₃	F	CN	CH, -SCH-COOCH,	0



R ¹	R ²	R³	R ⁴	R ⁵	x
-CCl ₃	СН3	F	CN	F	0
-CCl ₃	СН3	F .	CN	-0-CH ₂	0
-CCI ₃	СН3	F	CN	–o– ca- c≡ch ch²	0
-CCl ₃	СН3	а	CN	OCH ₃	O
-CCl ₃	СН3	а	CN	-S-C ₂ H ₅	0
-CCl ₃	СН3	F	NO ₂	OCH ₃	0
-CCl ₃	СН3	а	NO ₂	-O-CH2-C≡CH	0
-CCl ₃	сн3	а	CN	_O_CH_COOCH,	0
-CCl ₃	СН3	F	NO ₂	_O_CH_COOCH,	0
-CCI ₃	СН3	а	NO ₂	-o-ch-cooch,	0
-CC13	СН3	F	CN		0
-CCl ₃	CH ₃	F	CN	-O-CH ₂ -С _б Н ₅	0
CF ₃	СН3	F	CN	ОН	s

	R1	R²	R³	R ⁴	R ⁵	x
	CF ₃	СН3	F	CN	-O-i-C3H7	s
	CF3	CH ₃	F .	CN	-0-СH ₂ -СН=СН ₂	S
	CF ₃	CH ₃	F	CN	-O-CH2-C≡CH	S
	CF ₃	CH ₃	F	CN	СН, -0—СН-СООС,Н,	S
	CF ₃	CH ₃	F	CN	-0-CH ₂ -COOCH ₃	s
	CF ₃	СН3	F	CN	-S-CH ₃	s
	CF ₃	CH ₃	F	CN	-S-C ₂ H ₅	S
	CF ₃	СН3	F	CN	CH, -s—CH—COOCH,	S
	CF3	СН3	F	CN	F	s
5	CF ₃	CH3	F	CN	-0-CH ₂	S
,	CF ₃	СН3	F	CN	-o-a+c≡ah ah	S
	CF ₃	CH ₃	a	CN	OCH ₃	S
	CF ₃	CH ₃	a	CN	-S-C ₂ H ₅	s
	CF ₃	СН3	F	NO ₂	осн3	s.

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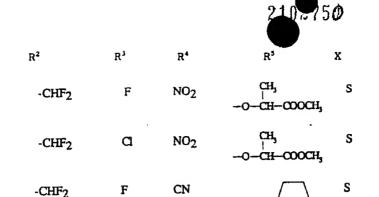


R1	R ²	R³	R ⁴	R ⁵	x
CF ₃	CH ₃	а	NO ₂	-O-CH2-C≡CH	S
CF ₃	СН3	α.	CN	—о—сн—соосн,	S
CF3	СН3	F	NO ₂	-0-сн-соосн, сн,	S
CF ₃	CH ₃	а	NO ₂	-0-CH-COOCH ²	S
CF ₃	CH ₃	F	CN	_o	S
CF ₃	CH ₃	F	CN	-O-CH ₂ -C ₆ H ₅	S
CF ₃	-CHF ₂	F	CN	ОН	S
CF ₃	-CHF ₂	F	CN	OCH ₃	S
CF ₃	-CHF ₂	F	CN	-O-CH ₂ -CH=CH ₂	s
CF ₃	-CHF ₂	F	CN	-O-CH2-C≡CH	s
CF ₃	-CHF ₂	F	CN	сн, о-сн-соос,н,	S

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R1	R ²	Ŕ³	R ⁴	R ⁵	X
CF ₃	-CHF ₂	F	CN	-0-CH ₂ -COOCH ₃	S
CF3	-CHF ₂	F	CN	-S-CH ₃	s
CF ₃	-CHF ₂	F	CN	-S-C ₂ H ₅	S
CF3	-CHF ₂	F	CN	CH, -SCH-COOCH,	s
CF ₃	-CHF ₂	F	CN	F	S
CF ₃	-CHF ₂	F	CN	-0-CH ₂ 0	S
CF ₃	-CHF2	F	CN	-o-ch-c≡ch Ch²	S
CF ₃	-CHF ₂	a.	CN	OCH ₃	s
CF3	-CHF ₂	а	CN	-S-C ₂ H ₅	s
CF ₃	-CHF ₂	F	NO ₂	OCH ₃	S
CF3	-CHF ₂	а	NO ₂	-O-CH2-C≡CH	S
CF ₃	-CHF ₂	а	CN	_0_CH_COOCH,	S

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If, for example, 4-methyl-3-trifluoromethyl-1,2,4-triazolin-5-one and 2,4,5-trifluorobenzonitrile are used as starting materials, the course of the reaction of process (a) according to the invention can be represented by the following equation:

If, for example, 1-(4-cyano-2,5-difluorophenyl)-4-methyl-

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R1

CF₃

CF₃

CF₃

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3-trifluoromethyl-1,2,4-triazolin-5-one and 3-Dutin-2-ol are used as starting materials, the course of the reaction of process (b) according to the invention can be represented by the following equation:

If, for example, 1-(4-cyano-2,5-difluorophenyl)-4-amino-3-trifluoromethyl-1,2,4-triazolin-5-one and sodium nitrite are used as starting materials, the course of the reaction of process (c) according to the invention can be represented by the following equation:

If, for example, 1-(4-cyano-2,5-difluorophenyl)-3-trifluoromethyl-(4H)-1,2,4-triazolin-5-one and chlorodifluoromethane are used as starting materials, the course of the reaction of process (d) according to the invention can be represented by the following equation:

Formula (provides a general definition the lH-triazolinones required as starting materials for carrying out process (a) according to the invention. In this formula (II), R¹, R² and X preferably and particularly preferably represent those radicals which have already been mentioned in connection with the description of the compounds of the formula (I) according to the invention as being preferred and particularly preferred for these substituents.

The lH-triazolinones of the formula (II) are known or can 10 be obtained analogously to known processes (compare, for US 4,477,459; DE 2,716,707; example, EP 399,294; 335-338 [1971]; US 3,780,052; J. Med. Chem. 14, DE 2,029,375). The compound 4-amino-3-trifluoromethyl-1H-15 1,2,4-triazolin-5-one was hitherto unknown and is also a subject of the invention. It is obtained when hydrazine hydrate is reacted first with diphenyl carbonate and subsequently with trifluoroacetic acid at temperatures between -20°C and +200°C (compare in this context also 20 the preparation examples).

Formula (III) provides a general definition of the halogenobenzene derivatives furthermore required as starting materials for carrying out process (a) according to the invention. In this formula (III), R³, R⁴ and R⁵ preferably and particularly preferably represent those radicals which have already been mentioned in connection with the description of the compounds of the formula (I) according to the invention as being preferred and

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particularly preferred for these substitutes. Hal preferably represents fluorine, chlorine or bromine, in particular fluorine or chlorine.

The halogenobenzene derivatives of the formula (III) have been disclosed or can be obtained in analogy to known processes (compare, for example, EP 191,181; EP 441,004; EP 431,373). The compound 5-chloro-2,4-difluorobenzo-nitrile was hitherto unknown and is also a subject of the invention. It is obtained when the known compound 2,4,5-trichlorobenzonitrile (compare, for example, EP 441,004) is reacted with potassium fluoride, if appropriate in the presence of a diluent such as, for example, tetramethylene sulphone, at temperatures between 100°C and 200°C (compare in this context also the Preparation Examples).

Formula (Ia) provides a general definition of the substituted triazolinones required as educts for carrying out process (b) according to the invention. In this formula (Ia), R¹, R², R³, R⁴ and X preferably and particularly preferably represent those radicals which have already been mentioned in connection with the description of the substances of the formula (I) according to the invention as being preferred and particularly preferred for these substituents. R⁵⁻¹ preferably represents fluorine, chlorine or bromine, in particular fluorine or chlorine.

The substituted triazolinones of the formula (Ia) are

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compounds according to the invention and can be obtained with the aid of processes (a), (c) and/or (d) according to the invention.

Formula (IV) provides a general definition of the nucleophiles furthermore required as educts for carrying out process (b) according to the invention. In this formula (IV), Z preferably represents oxygen or sulphur. R6-1 preferably and particularly preferably represents those radicals which have already been mentioned in connection with the description of the substances of the formula (I) according to the invention as being preferred and particularly preferred for the substituent R6 with the exception of the hydrogen radical. In the event that Z represents oxygen, R6-1 furthermore also preferably represents heterocyclyl, with a five- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms, in particular nitrogen, oxygen and/or sulphur, preferably being mentioned as heterocyclyl radical.

The nucleophiles of the formula (IV) are generally known compounds of organic chemistry.

Formula (Ib) provides a general definition of the substituted triazolinones required as educts for carrying out process (c) according to the invention. In this formula (Ib), R¹, R³, R⁴, R⁵ and X preferably and particularly preferably represent those radicals which have already been mentioned in connection with the description

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of the substances of the formula (I) according to the invention as being preferred and particularly preferred for these substituents. R^{2-1} preferably represents amino.

The substituted triazolinones of the formula (Ib) are compounds according to the invention and can be obtained with the aid of processes (a), (b) and/or (d) according to the invention.

Formula (Ic) provides a general definition of the substituted triazolinones required as educts for carrying out process (d) according to the invention. In this formula (Ic), R^1 , R^3 , R^4 , R^5 and X preferably and particularly preferably represent those radicals which have already been mentioned in connection with the description of the substances of the formula (I) according to the invention as being preferred and particularly preferred for these substituents. R^{2-2} preferably represents hydrogen.

The substituted triazolinones of the formula (Ic) are compounds according to the invention and can be obtained with the aid of processes (a), (b) and/or (c) according to the invention.

Formula (V) provides a general definition of the alkylating agents furthermore required as educts for carrying out process (d) according to the invention. In this formula (V), R^{2-3} preferably and particularly preferably represents those radicals which have already been

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mentioned in connection with the description of the substances of the formula (I) according to the invention as being preferred and particularly preferred for the substituent R2, with the exception of the radicals hydrogen, amino, cyano and alkylideneimino. E preferably represents a leaving radical which is customary in alkylating agents such as, for example, halogen, in particular chlorine, bromine or iodine, or in each case optionally substituted alkylsulphonyloxy, sulphonyloxy or arylsulphonyloxy such as, in particular, trifluoromethanesulphonyloxy, methanesulphonyloxy, ethoxysulphonyloxy methoxysulphonyloxy, p-toluenesulphonyloxy.

The alkylating agents of the formula (V) are generally known compounds of organic chemistry.

Suitable diluents for carrying out process (a) according to the invention are inert organic solvents. These include, in particular, aliphatic, alicyclic or aromatic, optionally halogenated hydrocarbons such as, for example, chlorobenzene, benzene, toluene, xylene, dichlorobenzene, petroleum ether, hexane, cyclohexane, dichloromethane, chloroform or carbon tetrachloride; ethers such as diethyl ether, diisopropyl ether, dioxane, tetrahydrofuran or ethylene glycol dimethyl ether or ethylene glycol diethyl ether; ketones such as acetone, butanone or methyl-isobutyl-ketone; nitriles such as acetonitrile, propionitrile or benzonitrile; amides such N, N-dimethylformamide, N, N-dimethylacetamide, as

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N-methylformanilide, N-methylpyrrolidone or hexamethylphosphoric triamide, or esters such as methyl acetate or ethyl acetate.

Process (a) according to the invention is preferably carried out in the presence of a suitable reaction auxiliary. Possible reaction auxiliaries are all customary inorganic or organic bases. These preferably include alkaline earth metal hydroxides or alkali metal hydroxides such as sodium hydroxide, calcium hydroxide, potassium hydroxide or else ammonium hydroxide, alkali metal carbonates such as sodium carbonate, potassium carbonate, potassium hydrogencarbonate, sodium hydrogencarbonate or ammonium carbonate, alkali metal acetates or alkaline earth metal acetates such as sodium acetate, potassium acetate, calcium acetate or ammonium acetate, and also tertiary amines such as trimethylamine, triethylamine, tributylamine, N.N-dimethylaniline, pyridine, N-methylpiperidine, piperidine, N, N-dimethylaminopyridine, diazabicyclooctane (DABCO), diazabicyclononene (DBN) or diazabicycloundecene (DBU).

When carrying out process (a) according to the invention, the reaction temperatures can be varied within a substantial range. In general, the process is carried out at temperatures between 0°C and +180°C, preferably at temperatures between +20°C and +120°C.

Process (a) according to the invention is conventionally carried out under atmospheric pressure. However, it is

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also possible to carry out the process under elevated or reduced pressure.

To carry out process (a) according to the invention, 1.0 to 3.0 mol, preferably 1.0 to 1.5 mol, of halogenobenzene derivative of the formula (III) and, if appropriate, 1.0 to 3.0 mol, preferably 1.0 to 1.5 mol, of base as reaction auxiliary are generally employed per mole of 1H-triazolinone of the formula (II). The reaction is carried out and the reaction products are worked up and isolated by known methods (compare in this context also the preparation examples).

Possible diluents for carrying out process (b) according to the invention are inert organic solvents. Preferably used solvents are those which have been listed in the description of process (a) according to the invention.

Process (b) according to the invention is preferably carried out in the presence of a suitable reaction auxiliary. Possible reaction auxiliaries are all customary inorganic or organic bases. These include, for example, the hydrides, hydroxides, amides, alcoholates, acetates, carbonates or hydrogencarbonates of alkaline earth metals or alkali metals such as, for example, sodium hydride, sodium amide, sodium methylate, sodium ethylate, potassium tert.-butylate, sodium hydroxide, potassium hydroxide, ammonium hydroxide, sodium acetate, potassium acetate, calcium acetate, ammonium acetate, sodium carbonate, potassium carbonate, potassium

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hydrogencarbonate, sodium hydrogencarbonate or ammonium carbonate, and also tertiary amines such as trimethylamine, triethylamine, N.N-dimethylaniline, pyridine, N-methylpiperidine, N.N-dimethylaminopyridine, diazabicyclooctane (DABCO), diazabicyclononene (DNB) or diazabicycloundecene (DBU).

When carrying out process (b) according to the invention, the reaction temperatures can be varied within a substantial range. In general, the process is carried out at temperatures between -20° C and $+150^{\circ}$ C, preferably at temperatures between 0° C and $+120^{\circ}$ C.

Process (b) according to the invention is conventionally carried out under atmospheric pressure. However, it is also possible to carry out the process under elevated or reduced pressure.

To carry out process (b) according to the invention, 1.0 to 3.0 mol, preferably 1.0 to 1.5 mol, of nucleophile of the formula (IV) and, if appropriate, 0.1 to 3.0 mol, preferably 1.0 to 1.5 mol, of base as reaction auxiliary are generally employed per mole of substituted triazolinone of the formula (Ia).

The reaction is carried out and the reaction products are worked up and isolated by known methods (compare in this context also the preparation examples).

25 Process (c) according to the invention is conventionally

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carried out in the presence of a suitable acid. Possible acids are, in particular, aqueous mineral acids. Dilute hydrochloric acid is particularly preferably used.

Suitable diluents for carrying out process (c) according to the invention are all diluents which are customary for such diazotisation reactions. It is particularly preferred to use a suitable excess of the aqueous mineral acids which have been employed as reagents, such as, for example, hydrochloric acid, simultaneously as the diluent.

When carrying out process (c) according to the invention, the reaction temperatures can be varied within a substantial range. In general, the process is carried out at temperatures between -20°C and +100°C, preferably at temperatures between -10°C and +80°C.

Process (c) according to the invention is conventionally carried out under atmospheric pressure. However, it is also possible to carry out the process under elevated or reduced pressure.

To carry out process (c) according to the invention, 1.0 to 3.0 mol, preferably 1.0 to 2.0 mol, of sodium nitrite and 1.0 to 10.0 mol, preferably 1.0 to 5.0 mol, of acid are generally employed per mole of substituted triazolinone of the formula (Ib).

25 The reaction is carried out and the reaction products are

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worked up and isolated by known methods (compare in this context also the preparation examples).

Possible diluents for carrying out process (d) according to the invention are inert organic solvents. These include, in particular, aliphatic, alicyclic or aromatic, optionally halogenated hydrocarbons such as, for example, benzine, benzene, toluene, xylene, chlorobenzene, dichlorobenzene, petroleum ether, hexane, cyclohexane, dichloromethane, chloroform, carbon tetrachloride; ethers such as diethyl ether, disopropyl ether, dioxane, tetrahydrofuran or ethylene glycol dimethyl ether or ethylene glycol diethyl ether; ketones such as acetone, butanone or methyl isobutyl ketone; nitriles such as acetonitrile, propionitrile or benzonitrile; amides such N, N-dimethylacetamide, N, N-dimethylformamide, N-methylformanilide, N-methylpyrrolidone or hexamethylphosphoric triamide; esters such as methyl acetate or ethyl acetate, or sulphoxides such dimethyl sulphoxide.

If appropriate, process (d) according to the invention can also be carried out in a two-phase system such as, for example, water/toluene or water/dichloromethane, if appropriate in the presence of a suitable phase transfer catalyst. Examples of such catalysts which may be mentioned are: tetrabutylammonium iodide, tetrabutylammonium bromide, tetrabutylammonium chloride, tributyl-methyl-phosphonium bromide, trimethyl-C₁₃/C₁₅-alkylammonium bromide, trimethyl-C₁₃/C₁₅-alkylammonium bromide,

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dibenzyl-dimethyl-ammoniummethylsulphate, dimethyl- C_{12}/C_{14} -alkyl-benzylammonium chloride, dimethyl- C_{12}/C_{14} -alkyl-benzylammonium bromide, tetrabutylammonium hydroxide, triethylbenzylammonium chloride, methyltrioctylammonium chloride, trimethylbenzylammonium chloride, 15-crown-5, 18-crown-6 or tris-[2-(2-methoxyethoxy)-ethyl]-amine.

Process (d) according to the invention is preferably carried out in the presence of a suitable reaction auxiliary. Suitable reaction auxiliaries are all customary inorganic or organic bases. These include, for example, the hydrides, hydroxides, amides, alcoholates, acetates, carbonates or hydrogencarbonates of alkaline earth metals or alkali metals such as, for example, sodium hydride, sodium amide, sodium methylate, sodium ethylate, potassium tert.-butylate, sodium hydroxide, potassium hydroxide, ammonium hydroxide, sodium acetate, potassium acetate, calcium acetate, ammonium acetate, carbonate, potassium carbonate, sodium hydrogencarbonate, sodium hydrogencarbonate or ammonium carbonate, and also tertiary amines such as trimethylamine, triethylamine, tributylamine, N,N-dimethylaniline, pyridine, N-methylpiperidine, N,N-dimethylaminopyridine, diazabicyclooctane (DABCO), diazabicyclononene (DBN) or diazabicycloundecene (DBU).

When carrying out process (d) according to the invention, the reaction temperatures can be varied within a substantial range. In general, the process is carried out at temperatures between -20°C and +150°C, preferably at

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temperatures between 0°C and +120°C.

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Process (d) according to the invention is conventionally carried out under atmospheric pressure. However, it is also possible to carry out the process under elevated or reduced pressure.

To carry out process (d) according to the invention, 1.0 to 3.0 mol, preferably 1.0 to 2.0 mol, of alkylating agent of the formula (V) and, if appropriate, 1.0 to 3.0 mol, preferably 1.0 to 2.0 mol, of base as reaction auxiliary are generally employed per mole of substituted triazolinone of the formula (Ic).

The reaction is carried out and the reaction products are worked up and isolated by known methods (compare in this context also the preparation examples).

The end products of the formula (I) are purified with the aid of conventional methods, for example by column chromatography or by recrystallisation.

They are characterised with the aid of the melting point or, in the case of compounds which do not crystallise, with the aid of proton nuclear resonance spectroscopy (1H NMR).

The active compounds according to the invention can be used as defoliants, desiccants, agents for destroying broad-leaved plants and, especially, as weed-killers. By

weeds, in broadest sense, there are to be inderstood all plants which grow in locations where they are undesired. Whether the substances according to the invention act as total or selective herbicides depends essentially on the amount used.

The active compounds according to the invention can be used, for example, in connection with the following plants:

Dicotyledon weeds of the genera: Sinapis, Lepidium,
Galium, Stellaria, Matricaria, Anthemis, Galinsoga,
Chenopodium, Urtica, Senecio, Amaranthus, Portulaca,
Xanthium, Convolvulus, Ipomoea, Polygonum, Sesbania,
Ambrosia, Cirsium, Carduus, Sonchus, Solanum, Rorippa,
Rotala, Lindernia, Lamium, Veronica, Abutilon, Emex,
Datura, Viola, Galeopsis, Papaver, Centaurea, Trifolium,
Ranunculus and Taraxacum.

Dicotyledon cultures of the genera: Gossypium, Glycine, Beta, Daucus, Phaseolus, Pisum, Solanum, Linum, Ipomoea, Vicia, Nicotiana, Lycopersicon, Arachis, Brassica, Lactuca, Cucumis and Cucurbita.

Monocotyledon weeds of the genera: Echinochloa, Setaria, Panicum, Digitaria, Phleum, Poa, Festuca, Eleusine, Brachiaria, Lolium, Bromus, Avena, Cyperus, Sorg hum, Agropyron, Cynodon, Monochoria, Fimbristylis, Sagittaria, Eleocharis, Scirpus, Paspalum, Ischaemum, Sphenoclea, Dactyloctenium, Agrostis, Alopecurus and Apera.

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Monocotyle Cultures of the genera: 2a, Zea, Triticum, Hordeum, Avena, Secale, Sorghum, Panicum, Saccharum, Ananas, Asparagus and Allium.

However, the use of the active compounds according to the invention is in no way restricted to these genera, but also extends in the same manner to other plants.

The compounds are suitable, depending on the concentration, for the total combating of weeds, for example on industrial terrain and rail tracks, and on paths and squares with or without tree plantings. Equally, the compounds can be employed for combating weeds in perennial cultures, for example afforestations, decorative tree plantings, orchards, vineyards, citrus groves, nut orchards, banana plantations, coffee plantations, tea plantations, rubber plantations, oil palm plantations, cocoa plantations, soft fruit plantings and hopfields, and for the selective combating of weeds in annual cultures.

The active compounds according to the invention can also be used particularly successfully for combating mono- and dicotyledon weeds.

The active compounds are furthermore suitable for combating animal pests, preferably arthropods and nematodes, in particular insects and arachnids, encountered in agriculture, in forestry, in the protection of stored products and of materials, and in the hygiene field. They

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are active against normally sensitive and resistant species and against all or some stages of development.

The abovementioned pests include:

From the order of the Isopoda, for example, Oniscus asellus, Armadillidium vulgare and Porcellio scaber; from the order of the Diplopoda, for example, Blaniulus guttulatus; from the order of the Chilopoda, for example, Geophilus carpophagus and Scutigera spec.;

from the order of the Symphyla, for example, Scutigerella immaculata;

from the order of the Thysanura, for example, Lepisma saccharina;

from the order of the Collembola, for example, Onychiurus

from the order of the Orthoptera, for example, Blatta orientalis, Periplaneta americana, Leucophaea maderae, Blattella germanica, Acheta domesticus, Gryllotalpa spp., Locusta migratoria migratorioides, Melanoplus

differentialis and Schistocerca gregaria; from the order of the Dermaptera, for example, Forficula auricularia;

from the order of the Isoptera, for example, Reticulitermes spp.;

from the order of the Anoplura, for example, Phylloxera vastatrix, Pemphigus spp., Pediculus humanus corporis, Haematopinus spp. and Linognathus spp.;

from the order of the Mallophaga, for example,

Trichodectes spp. and Damalinea spp.; from the order of the Thysanoptera, for example, Hercinothrips femoralis and Thrips tabac1; from the order of the Heteroptera, for example, Eurigaster spp., Dysdercus intermedius, Piesma quadrata, 5 Cimex lectularius, Rhodnius prolixus and Triatoma spp.; from the order of the Homoptera, for example, Aleurodes brassicae, Bemisia tabaci, Trialeurodes vaporariorum, Aphis gossypii, Brevicoryne brassicae, Cryptomyzus ribis, lanigerum, Eriosoma fabae, Doralis pomi, 10 Hyalopterus arundinis, Macrosiphum avenae, Myzus spp., Phorodon humuli, Rhopalosiphum padi, Empoasca spp., Euscelis bilobatus, Nephotettix cincticeps, Lecanium striatellus, Laodelphax oleae, Saissetia corni, Nilaparvata lugens, Aonidiella aurantii, Aspidiotus 15 hederae, Pseudococcus spp. and Psylla spp.; from the order of the Lepidoptera, for example, Pectinophora gossypiella, Bupalus piniarius, Cheimatobia brumata, Lithocolletis blancardella, Hyponomeuta padella, Plutella maculipennis, Malacosoma neustria, Euproctis 20 chrysorrhoea, Lymantria spp., Bucculatrix thurberiella, Phyllocnistis citrella, Agrotis spp., Euxoa spp., Feltia spp., Earias insulana, Heliothis spp., Laphygma exigua, Mamestra brassicae, Panolis flammea, Prodenia litura, Spodoptera spp., Trichoplusia ni, Carpocapsa pomonella, 25 Pieris spp., Chilo spp., Pyrausta nubilalis, Ephestia kuehniella, Galleria mellonella, Tineola bisselliella, pseudospretella, Hofmannophila pellionella, Tinea Choristoneura reticulana, podana, Capua Cacoecia fumiferana, Clysia ambiguella, Homona magnanima and 30

Tortrix viridana;

from the order of the Coleoptera, for example, Anobium punctatum, Rhizopertha dominica, Bruchidius obtectus, Acanthoscelides obtectus, Hylotrupes bajulus, Agelastica 5 alni, Leptinotarsa decemlineata, Phaedon cochleariae, Diabrotica spp., Psylliodes chrysocephala, Epilachna varivestis, Atomaria spp., Oryzaephilus surinamensis, Anthonomus spp., Sitophilus spp., Otiorrhynchus sulcatus, Cosmopolites sordidus, Ceuthorrhynchus assimilis, Hypera 10 postica, Dermestes spp., Trogoderma spp., Anthrenus spp., Attagenus spp., Lyctus spp., Meligethes aeneus, Ptinus spp., Niptus hololeucus, Gibbium psylloides, Tribolium spp., Tenebrio molitor, Agriotes spp., Conoderus spp., Melolontha melolontha, Amphimallon solstitialis and 15 Costelytra zealandica;

from the order of the Hymenoptera, for example, Diprion spp., Hoplocampa spp., Lasius spp., Monomorium pharaonis and Vespa spp.;

from the order of the Diptera, for example, Aedes spp.,

20 Anopheles spp., Culex spp., Drosophila melanogaster,

Musca spp., Fannia spp., Calliphora erythrocephala,

Lucilia spp., Chrysomyia spp., Cuterebra spp.,

Gastrophilus spp., Hyppobosca spp., Stomoxys spp.,

Cestrus spp., Hypoderma spp., Tabanus spp., Tannia spp.,

Bibio hortulanus, Oscinella frit, Phorbia spp., Pegomyia

hyoscyami, Ceratitis capitata, Dacus oleae and Tipula paludosa;

from the order of the Siphonaptera, for example, Kenopsylla cheopis and Ceratophyllus spp.;

30 from the order of the Arachnida, for example, Scorpio

maurus and Latrodectus mactans;

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from the order of the Acarina, for example, Acarus siro, Argas spp., Ornithodoros spp., Dermanyssus gallinae, Eriophyes ribis, Phyllocoptruta oleivora, Boophilus spp., Rhipicephalus spp., Amblyomma spp., Hyalomma spp., Ixodes spp., Psoroptes spp., Chorioptes spp., Sarcoptes spp., Tarsonemus spp., Bryobia praetiosa, Panonychus spp. and Tetranychus spp..

The active compounds according to the invention are distinguished by a powerful insecticidal and acaricidal activity. They can be used particularly successfully for combating the greenhouse red spider mite (Tetranychus urticae). Besides, the active compounds have, in particular, leaf-acting insecticidal properties.

Depending on their particular physical and/or chemical properties, the active compounds can be converted to the customary formulations, such as solutions, emulsions, suspensions, powders, foams, pastes, granules, aerosols, natural and synthetic materials impregnated with active compound, very fine capsules in polymeric substances and in coating compositions for seed, furthermore in formulations used with burning equipment, such as fumigating cartridges, fumigating cans, fumigating coils and the like, as well as ULV cold mist and warm mist formulations.

These formulations are produced in a known manner, for example by mixing the active compounds with extenders,

that is, liquid solvents, liquefied gases under pressure, and/or solid carriers, optionally with the use of surface-active agents, that is emulsifying agents and/or dispersing agents and/or foam-forming agents. In the case of the use of water as an extender, organic solvents can, for example, also be used as auxiliary solvents. As liquid solvents, there are suitable in the main: aromatics, such as xylene, toluene or alkylnaphthalenes, chlorinated aromatics or chlorinated aliphatic hydrocarbons, such as chlorobenzenes, chloroethylenes or methylene chloride, aliphatic hydrocarbons, such as cyclohexane or paraffins, for example mineral oil fractions, alcohols, such as butanol or glycol as well as their ethers and esters, ketones, such as acetone, methyl ethyl ketone, methyl isobutyl ketone or cyclohexanone, strongly polar solvents, such as dimethylformamide and dimethyl sulphoxide, as well as water; by liquefied gaseous extenders or carriers are meant liquids which are gaseous at ambient temperature and under atmospheric pressure, for example aerosol propellants, such as halogenated hydrocarbons as well as butane, propane, nitrogen and carbon dioxide; as solid carriers there are suitable: for example ground natural minerals, such as kaolins, clays, talc, chalk, quartz, attapulgite, montmorillonite or diatomaceous earth, and ground synthetic minerals, such as highly disperse silica, alumina and silicates; as solid carriers for granules there are suitable: for example crushed and fractionated natural rocks such as calcite, marble, pumice, sepiolite and dolomite, as well as synthetic granules of inorganic and

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organic means, and granules of organic materal such as sawdust, coconut shells, maize cobs and tobacco stalks; as emulsifying and/or foam-forming agents there are suitable: for example non-ionic and anionic emulsifiers, such as polyoxyethylene fatty acid esters, polyoxyethylene fatty alcohol ethers, for example alkylaryl polyglycol ethers, alkylsulphonates, alkyl sulphates, arylsulphonates as well as albumen hydrolysis products; as dispersing agents there are suitable: for example lignin-sulphite waste liquors and methylcellulose.

Adhesives such as carboxymethylcellulose and natural and synthetic polymers in the form of powders, granules or latices, such as gum arabic, polyvinyl alcohol and polyvinyl acetate, as well as natural phospholipids, such as cephalins and lecithins, and synthetic phospholipids, can be used in the formulations. Other additives can be mineral and vegetable oils.

It is possible to use colorants such as inorganic pigments, for example iron oxide, titanium oxide and Prussian Blue, and organic dyestuffs, such as alizarin dyestuffs, azo dyestuffs and metal phthalocyanine dyestuffs, and trace nutrients such as salts of iron, manganese, boron, copper, cobalt, molybdenum and zinc.

The formulations in general contain between 0.1 and 95 per cent by weight of active compound, preferably between 0.5 and 90%.

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When used as herbicides, the active compounds according to the invention, as such or in the form of their formulations, can also be used as mixtures with known herbicides, when used as herbicides, finished formulations or tank mixes being possible.

Suitable herbicides for the mixtures are known herbicides, for example anilides such as, for example, diflufenican and propanil; arylcarboxylic acids such as, for example, dichloropicolinic acid, dicamba or picloram; aryloxyalkanoic acids such as, for example, 2,4-D, 2,4-2,4-DP, fluroxypur, MCPA, MCPP and triclopyr: aryloxy-phenoxy-alkanoic esters such as, for example, diclofop-methyl, fenoxaprop-ethyl, fluazifop-butyl, haloxyfop-methyl and quizalofop-ethyl; azinones such as, for example, chloridazon and norflurazon; carbamates such as, for example, chlorpropham, desmedipham, phenmedipham and propham; chloroacetanilides such as, for example, alachlor, acetochlor, butachlor, metazachlor, metolachlor, pretilachlor and propachlor; dinitroanilines such as, for example, oryzalin, pendimethalin and trifluralin; diphenyl ethers such as, for example, acifluorfen. bifenox, fluoroglycofen, fomesafen, halosafen, lactofen and oxyfluorfen; ureas such as, for example, chlortoluron, diuron, fluometuron, isoproturon, linuron and methabenzthiazuron; hydroxylamines such as, for example, alloxydim, clethodim, cycloxydim, sethoxydim and tralkoxydim; imidazolinones such as, for example, imazethapyr, imazamethabenz, imazapyr and imazaquin; nitriles such as, for example, bromoxynil, dichlobenil

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oxvacetamides such as. mefenacet; sulphonylureas such as, for example, amidosulfuron, bensulfuron-methyl, chlorimuron-ethyl, chlorsulfuron, cinosulfuron, metsulfuron-methyl, nicosulfuron, pyrazosulfuron-ethyl, thifensulfuronprimisulfuron, methyl triasulfuron and tribenuron-methyl; thiocarbamates such as, for example, butylate, cycloate, di-allate, EPTC, esprocarb, molinate, prosulfocarb, thiobencarb and tri-allate; triazines such as, for example, atrazine, cyanazine, simazine, simetryn, terbutryn and terbutylazine; triazinones such as, for example, hexazinone, metamitron and metribuzin; others such as, for example, aminotriazole, benfuresate, bentazone, cinmethylin. clomazone, clopyralid, difenzoquat, dithiopyr, ethofumesate, fluorochloridone, glufosinate, glyphosate, isoxaben, pyridate, quinchlorac, quinmerac, sulphosate and tridiphane.

Mixtures with other known active compounds, such as fungicides, insecticides, acaricides, nematicides, bird repellants, plant nutrients and agents which improve soil structure, are also possible.

When used as herbicides, the active compounds can be used as such, in the form of their formulations or in the use forms prepared therefrom by further dilution, such as ready-to-use solutions, suspensions, emulsions, powders, pastes and granules. They are used in the customary manner, for example by watering, spraying, atomizing or scattering.

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When used as herbicides, the active compounds according to the invention can be applied either before or after emergence of the plants.

They can also be incorporated into the soil-before sowing.

When used as herbicides, the amount of active compound used can vary within a substantial range. It depends essentially on the nature of the desired effect. In general, the amounts used are between 0.01 and 10 kg of active compound per hectare of soil surface, preferably between 0.05 and 5 kg per hectare.

When used as insecticides and acaricides, the active compounds according to the invention can also be present in their commercially available formulations and in the use forms, prepared from these formulations, as a mixture with other active compounds, such as insecticides, attractants, sterilising agents, acaricides, nematicides, fungicides, growth-regulating substances or herbicides. The insecticides include, for example, phosphates, carbamates, carboxylates, chlorinated hydrocarbons, phenylureas and substances produced by microorganisms.

When used as insecticides and acaricides, the active compounds according to the invention can furthermore be present in their commercially available formulations and in the use forms, prepared from these formulations, as a mixture with synergistic agents. Synergistic agents are

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compounds which increase the action of the active compounds, without it being necessary for the synergistic agent added to be active itself.

The active compound content of the use forms prepared from the commercially available formulations can vary within wide limits. The active compound concentration of the use forms can be from 0.0000001 to 95 per cent by weight of active compound, preferably between 0.0001 and 1 per cent by weight.

When used as insecticides and acaricides, the compounds are employed in a customary manner appropriate for the use forms.

The preparation and the use of the active compounds according to the invention can be seen from the Examples which follow.

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Preparation Examples:

Example 1:

(Process a)

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5.3 g (0.038 mol) of potassium carbonate is added at room temperature to 5.3 g (0.032 mol) of 4-methyl-3-trifluoromethyl-1,2,4-triazolin-5-one (compare, for example, US 3,780,052) and 5.5 g (0.032 mol) of 5-chloro-2,4difluorobenzonitrile in 100 ml of dimethyl sulphoxide, and the mixture is subsequently heated for 36 hours at 100°C. For work-up, the cooled reaction mixture is poured into water, the pH is brought to 2 using dilute hydrochloric acid, and the mixture is extracted several times using dichloromethane. The combined organic phases are dried over sodium sulphate and concentrated in vacuo. The residue chromatographed over silica ael (eluent:dichloromethane).

1.8 g (18 % of theory) of 1-(2-chloro-4-cyano-5-fluoro-phenyl)-4-methyl-3-trifluoromethyl-1,2,4-triazolin-5-one



of melting point 105°C are obtained.

Example 2:

Process (b)

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0.6 g (0.014 mol) of sodium hydride (60 % in mineral oil) is added with stirring at room temperature to 1.0 g (0.014 mol) of 3-butin-2-ol in 50 ml of acetonitrile, the mixture is stirred for 15 minutes at room temperature, 2.9 g (0.01 mol) of 1-(2,5-difluoro-4-cyano-phenyl)-4-methyl-3-trifluoromethyl-1,2,4-triazolin-5-one are then added, and the mixture is subsequently stirred for a further 2 hours at room temperature. For work-up, the reaction mixture is concentrated in vacuo, the residue is partitioned between dichloromethane and water, and the organic phase is dried over sodium sulphate and freed from solvent in vacuo.

1.8 g (54 % of theory) of 1-(2-fluoro-4-cyano-5-but-1-in-3-yl-oxy-phenyl)-4-methyl-3-trifluoromethyl-1,2,4-triazolin-5-one of melting point 41°C are obtained.

Example 3:

Process (a)

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1.7 g (0.012 mol) of potassium carbonate are added at room temperature to 1.7 g (0.01 mol) of 4-amino-3-trifluoromethyl-1,2,4-triazolin-5-one and 1.6 g (0.01 mol) of 2,4,5-trifluorobenzonitrile (compare, for example, EP 191,181) in 30 ml of dimethyl sulphoxide, and the mixture is subsequently stirred for a further 14 hours at room temperature. For work-up, the reaction mixture is transferred into water, the pH is brought to 2 using dilute hydrochloric acid, and the mixture is extracted several times using dichloromethane. The combined organic phases are dried over sodium sulphate and concentrated in vacuo, and the residue is stirred with water, filtered off with suction and dried.

2.6 g (87 % of theory) of 1-(2,5-difluoro-4-cyano-phenyl)-4-amino-3-trifluoromethyl-1,2,4-triazolin-5-one of melting point 141°C are obtained.

Example 4:

Process (C)

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A saturated aqueous solution of 1.4 g (0.02 mol) of sodium nitrite is added at -5°C to 0°C in the course of 15 minutes with stirring to 3.0 g (0.01 mol) of 1-(2,5-difluoro-4-cyano-phenyl)-4-amino-3-trifluoromethyl-1,2,4-triazolin-5-one in 40 ml of 10 % strength hydrochloric acid, the cold bath is subsequently removed, the mixture is stirred for 1 hour at room temperature and is then again cooled to -5°C to 0°C and filtered, and the residue is washed with water and dried.

1.8 g (62 % of theory) of 1-(2,5-difluoro-4-cyano-phenyl)-3-trifluoromethyl-1,2,4-triazolin-5-one of melting point 51°C are obtained.

Example 5:

Process (d)

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15 g (0.17 mol) of chlorodifluoromethane are passed at 0°C to 10°C in the course of 5 hours into a suspension of 2.5 g (0.009 mol) of 1-(2,5-difluoro-4-cyanophenyl)-3-trifluoromethyl-4H-1,2,4-triazolin-5-one, 1.0 g(0.017 mol) of potassium hydroxide and 0.25 g of tetrabutylammonium bromide in 50 ml of tetrahydrofuran, and, during this time, the consumption of base is compensated for after 1, 2 and 3 hours in each case by adding further 1.0 g portions (0.017 mol) of potassium hydroxide. For work-up, the reaction mixture is poured into water and extracted several times using ethyl acetate, the combined organic phases are dried over sodium sulphate, and the solvent is subsequently removed in vacuo. The residue is chromatographed over silica gel (eluent: dichloromethane).

2.2 g (75 % of theory) of 1-(2,5-difluoro-4-cyanophenyl)-3-trifluoromethyl-4-difluoromethyl-1,2,4-triazolin-5-one of melting point 68°C are obtained.

Preparation of the starting compounds:

Example II-1:

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2782 g (13 mol) of diphenyl carbonate are added in portions in the course of 2 hours with stirring and icecooling to 1300 g (26 mol) of hydrazine hydrate in such a way that the temperature of the reaction mixture does not rise above 30°C, the mixture is subsequently stirred for 2 hours at 80°C and then cooled again, and 3164 g (26 mol) of trifluoroacetic acid are added, also in portions. The mixture is then stirred for another 2 hours at 80°C, and water is subsequently distilled off until the residue has reached a temperature of 180°C. When cooled, 1100 g (16.2 mol) of aqueous ammonia (25 % strength) are added, and the mixture is heated for 3 hours at reflux temperature. For work-up, all volatile components are distilled off under gradually reduced pressure (down to 20 mbar) until the residue has reached a temperature of 180°C, and the residue is recrystallised from 2000 ml of water, filtered off with suction and dried.

702 g (32 % of theory) of 3-trifluoromethyl-4-amino-1H-1,2,4-triazolin-5-one of melting point 163°C are obtained.

Example III-1:

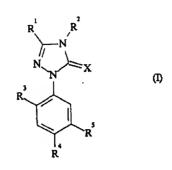
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220 g (1.06 mol) of 2,4,5-trichlorobenzonitrile (compare, for example, EP 441,004) are added with stirring at room temperature to 250 g (4.31 mol) of potassium fluoride in 400 ml of distilled tetramethylene sulphone, and the mixture is subsequently stirred for 10 hours at 195°C to 200°C. For work-up, the mixture is cooled, 500 ml of water are added, and the mixture is subjected to steam distillation. The organic portion is taken up in dichloromethane and the mixture is dried over sodium sulphate, concentrated in vacuo and distilled.

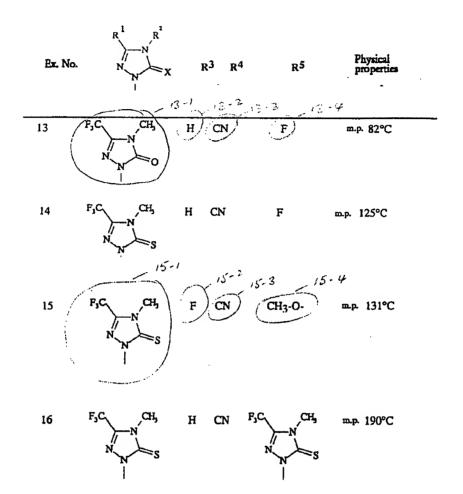
108 g (58 % of theory) of 2,4-difluoro-5-chlorobenzo-nitrile of boiling point $105-107^{\circ}C$ at 30 mbar and of melting point $48-50^{\circ}C$ are obtained.

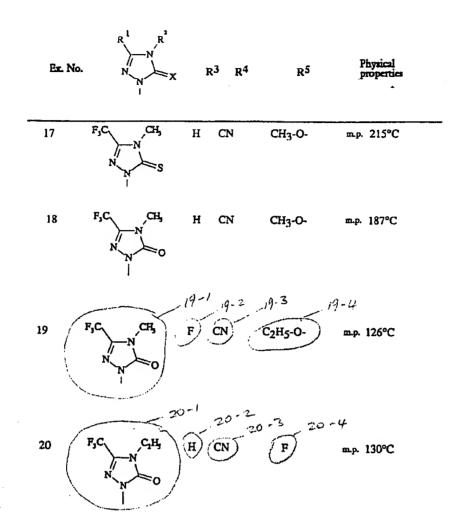
The following substituted triazolinones of the general formula (I) are obtained in a corresponding manner and following the general information on the preparation:



Ex. No.	R ¹ R ²	R ³ R	t ⁴ R ⁵	Physical properties
6	F,C,C,H,	F C	EN F	1 _{H NMR} *): 1.45 ⁻ -1.55; 4.22-4.3; 7.58-7.62
7	F ₃ C CH ₃	F (ON H	m.p. 99°C
8	F ₃ C CH ₃	CI N	то ₂ н	m.p. 110°C

 $\gamma)$ Н ш.р. 108°С CI CN 9 m.p. 103°C 11 12





Ex. No.	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	R ³ R ⁴	R ^S	Physical properties
21	F ₃ C C ₂ H ₃	H CN F ₃ C	C,H,	m.p. 138°C
22	F ₃ C C ₂ H ₃	F CN	F	m.p. 68°C
23	F ₃ C C ₂ H ₃	CI CN	Н	m.p. 145°C
24	F ₃ C C ₂ H ₃	F CN	н	≖.р. 204°С



Ex. No.	R ¹ R ²	R ²	8 R ⁴	R ⁵	Physical properties
25	F ₃ C C ₂ H ₃	F	CN	CH, O-CH-C≡CH	¹ H NMR*): 1.75-1.78; 2.6; 3.9-4.0
26	F ₃ C C ₂ H ₃	F	CN	СН3-О-	m.р. 133-135°С
27	F ₃ C C ₃ H ₃	F	CN	-NH-CH ₃	m.p. 143°C
28	F ₃ C N=C CH ₃	F	CN	F	m.p. 148°C

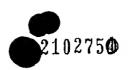
Ex. No	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	I	₹3 R	4 _R 5	Physical properties
29	F ₃ C CH ₃	F	CN	F	m.p. 74°C
30	F ₃ C CH ₃	F	CN	–o–ਕਮ-c≡ਕਮ ਖ਼ਿਸ਼ੇ	m.p. 116°C
31	F ₃ C n-C ₄ H ₂	F	CN	F	¹ H NMR [*]): 1.38-1.5; 1.73-1.83; 3.82-3.88
32	F ₂ CH, CH,	F	CN	F	ш.р. 177°С

Ex. No.	R ¹ R ² X	R ³ R ⁴	R ⁵	Physical properties
33	F ₃ C C ₂ H ₃	F NO ₂	F ₃ C C ₂ H ₃	m.p. 177°C
34	F ₃ C C ₂ H ₃	F CN -(0	O-CH ₂ -CH ₂) ₂ -OCH	3.48; 3.55-3.6; 3.9-3.97
35	F ₃ C C ₂ H ₃	F CN	-O-С ₂ Н ₅	¹ H NMR ^{*)} : 1.4-1.46; 1.5- 1.55; 3.9-3.98; 4.14-4.2
36	F ₃ C C ₂ H ₃	F CN	-0-i-C ₃ H ₇	¹ H NMR ^{*)} : 3.9-3.98; 4.6- 4.68; 7.2-7.23; 7.42-7.45

Ex. No.	R ¹ R ² X	R	3 R ⁴	R ⁵	Physical properties
37	F ₅ C, n-C ₆ H ₅	F	CN	—o—aн-c≡an	¹ H NMR [*]): 1.72-1.8; 3.8-3.87; 7.45-7.5
38	F ₃ C N O	F	CN	F	m.p. 90°C
39	F ₃ C N O	F	NO ₂	F	m-b∙ ∂∂₀C
40	F ₃ C N N O	F	CN	—0—Gi-C≡Gi Gi	m.p. 95°C

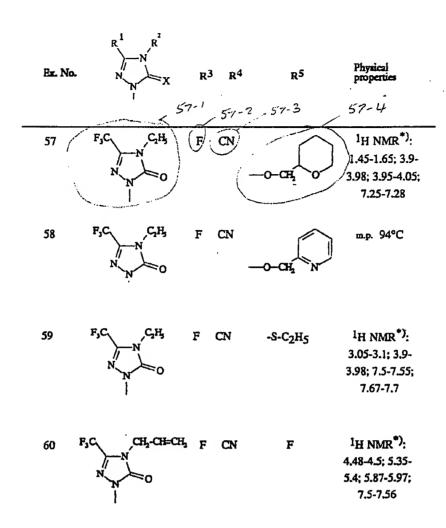
Ex. 1	√o.	R ¹	X X	R	3 R4	R ⁵	Physical properties
41	F ₃ C	CH ₂ -N	a a	F	CN	F	¹ H NMR [*]): 1.75-1.8; 2.08- 2.18; 3.85-3.92; 7.03-7.18
42	F ₃ C	CH ₂	a Ca	F	CN	—0— ai -c≡ai	¹ H NMR ^{*)} : 1.75-1.8; 4.33- 4.42; 4.9-4.98; 7.45-7.5
43	F30	N N	сн, e	F	CN	-O-CH ₂ -Si(CH ₃) ₃	m.p. 101°C
44	F ₃ (C N N	C'H'	F	CN	-O-CH ₂ -CH=CH ₂	m.p. 76°C

Ex. No.	R ¹ , R ²	R3	R4	R ⁵	Physical properties
45	F,C C,H,	F CN	-O-(CH ₂);	₂ -O-i-C ₃ H ₇	¹ H NMR ^{*)} : 1.18-1.22; 1.4- 1.45; 3.8-3.85; 4.22-4.25
46	F ₃ C C ₂ F ₃	F CN	-O-(CH ₂) ₂ -	СН(СН3)≖СН	1H NMR*): 1.85; 3.9- 3.98; 4.15- 4.2; 7.2-7.23
47	F ₃ C C ₂ H ₃	F CN	-O-CH(CH ₂	3)-CH ₂ -OCH	3.4; 3.9-3.98; 7.1-7.13; 7.38-7.42
48	F ₃ C C ₂ H ₃	α (en	F	m.p. 121°C



Ex. No.	R ¹ R ² X	R	3 R4	R ⁵	Physical properties
49	F ₃ C C ₂ H ₃	F	CN	-o-(och,	m.p. 154°C
50	F ₃ C C ₂ H ₃	F	CN	-N(CH ₃) ₂	¹ H NMR ^{*)} : 3.17; 3.9-3.98; 7.1-7.13; 7.38-7.42
51	F ₃ C C ₁ H ₃	а	CN	—o—aн-c≡aн	¹ H NMR ^{*)} : 1.75-1.8; 3.9- 3.98; 4.9-5.0; 7.35; 7.75
52	F,C CH,	а	CN	-O-CH3	mp 133°C

Ex. No.	R ¹ R ² X	R	3 R	4 _R 5	Physical properties
53	F,C CH,	F	CN	-O-n-C3H7	m.p. 71°C
54	F ₃ C , C ₂ H,	F	CN	—0-CH₂-C≡CH	¹ H NMR [*]): 2.53; 3.9-3.98; 4.85; 7.4-7.42
55	F ₃ C C ₂ H ₃	F	CN	-O-(CH ₂) ₂ -S-C ₂ H ₅	¹ H NMR ^{*)} : 2.67-2.78; 3.9- 3.98; 4.22-4.3; 7.23-7.25
56	F ₃ C , C ₂ H ₃	СІ	CN	CI CI	ш.р. 97°С



Ex. No.	R R X	R ³	R ⁴	R ⁵	Physical properties
61	F ₃ C CH ₃	F	CN 2	-O-n-C ₃ H ₇	mp.33°C
62	F ₃ C CH ₂ CH=CH ₂	F	(8)	CH3 CH3	¹ H-NMR: 1,75-1,78; 4,45-4,48; 7,45-7,50.
63	F ₃ C	ቴ	CX	-NH-CH₂CH=CH₂	¹ H-NMR: 1,40-1,45; 3,85-3,90; 6,83-6,86.
64	F ₃ C	F	3	-0-CH ₂	mp.101°C
64	F ₃ C	F	CN	_о_сн ₂ о	¹ H-NMR: 1,40-1,45; 4,08-4,15; 7,45-7,48.
64	F ₃ C	F	CN	—о—сн ₂ —	mp. 91°C
67	F ₃ C	F	CN	-0-CH(CH ₂ OC ₂ H ₅) ₂	¹ H-NMR: 3,53 - 3,60; 3,90-3,98; 4,55-4,60.

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Ex. No.	R R R	R ³	R ⁴	R ⁵	Physical properties
68	F ₃ C C ₂ H ₅	F	CN	-осн ₂ сн ₂ сн сн ₃	mp.81°C
69	F₂CH CH₃	F	CN	—CH—C≣CH CH₃	¹ H-NMR: 2,60;4,90- 4,98; 7,45-7,50.
70	F ₂ CH N C ₂ H ₅	F	CN	F	mp.161°C
71	F ₃ C	F	CN	—о—сн ₂ —△	mp.96°C
72	F ₂ CH CH ₃	F	СИ	ocHc≡ch	mp.176°C
73	F ₃ C	F	СИ	-O-(CH ₂ CH ₂ O) ₅ CH ₃	¹ H-NMR: 3,52-3,56; 3,60-3,70; 4,75-4,78.
74	F ₃ C	F	CN	-оվсн _у сн _у о _й сн _у сн ь сн _у	¹ H-NMR: 3,60-3,65; 3,88-3,96; 5,85-6,00.



Ex. No.	R R R	R ³	R ⁴	R ⁵	Physical properties
	R R R	F	CN	-O-CH ₂ CH=CHCH ₃	mp.117°C
75	F ₃ C	r	CIV		
76	F ₃ C	F	CN	—оснсн=сн ₂ сн ₃	mp. 47°C
78	F ₃ C	F	СИ	-0-CH ₂ CH ₂ N(CH ₃) ₂	5,82-5,95.
78	F ₃ C C ₂ H ₅	F	CN	-o-ch₂chc₂h ch₃	
79	F ₃ C N C ₂ H ₅	F	CN	-o-сңсңсң	mp.87°C
78	F ₃ C	F	CN	-0-CHC ₂ H ₅ CH ₃	¹ H-NMR: 3,90-3,98; 4,38-4,45; 7,43-7,46.
81	F ₃ C N C ₂ F	1 ₅ F	CN	-0-сң,сң,сң(с	1 ₃) ₂ mp. 75°C

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Ex. No.	R	N N X	R ³	R ⁴	R ⁵	Physical properties
82	F	F ₃ C N C ₂ H ₅	F	CN	-O-CH₂C(CH₃)₃	mp.117°C
83		F ₃ C	F	CN	-0-СН ₂ СI	mp.141°C
84		F ₃ C	F	CN	-O-CH ₂ CH ₂ -N	mp. 143°C
85		F ₃ C	F	CN	-о-Снсоосун сн,	¹ H-NMR: 3,85-3,92; 4,16-4,26; 4,70-4,76.
8	6	F ₃ C	F	CN	—o-¢нсн v(сн _з)	2,32; 3,90- 3,98; 4,53-4,60.
8	37	F ₃ C N C ₂ H ₅	F	CN	-осн,сн,осн,сн,исн	mp.65°C

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Ex. No.	R ¹	R³	R ⁴	R ⁵	Physical Properties
88	F ₃ C	F	CN	NHCHC₂H₅ I CH₃	mp. 91°C
89	F ₃ C	F	CN	—NH−CH(CH₃)₂	mp.100°C
90	F ₃ C N C ₂ H ₅	F	CN	NHC ₆ H₁₃n	mp.86°C
91	F ₃ C	F	CN	-NH-(H)	mp. 126°C
92	F ₃ C N C ₂ H ₅	F	NO ₂	F	mp. 81°C
93	F ₃ C	F	CN	-NHCH ₂ CH ₂ OCH ₃	mp. 57°C

Ex. No.	R R X	R³	R ⁴	R ⁵	Physical properties
94	F ₂ CH C ₂ H ₅	F	CN	F	mp.117°C -
95	F ₂ CH C ₂ H ₅	F	C Z	—o−chc <u>≕</u> ch ch³	mp.96°C
96	F₂CH C₂H₅	F	CN	O-CH₂C≡CH	⁴ H-NMR: 2,62-2,64; 3,95-4,02; 4,85.
97	F ₂ CH C ₂ H ₅	F	CN	осн сн₂осн, сн₃	mp.78°C
98	F ₂ CH CH ₃	F	CN	оснсң ₂ осң, сң,	¹ H-NMR: 1,28-1,30; 3,40; 3,50; 4,55-4,65.
99	F ₂ CH C ₂ H ₅	F	CN	—о—снсн _г осн _з сн _з	mp. 90°C

Ex. No.	R R R X	R ³	R ⁴	R ⁵	Physical properties
100	F ₂ CH C ₂ H ₅	F	CN	-O-CHC≡CH CH₃	mp_134°C
101	F ₂ CH CH ₃	F	CN	-O-CH(CH₃)₂	mp.135°C
102	F ₅ C ₂ CH ₃	F	СИ	F	mp. 96°C
103	F ₅ C ₂ CH ₃	F	CN	—O-ÇHC≣CH CH₃	mp.115°C
104	F ₅ C ₂ CH ₃	F	CN	OCH²C≣CH	
105	F ₂ CHCF ₂ CH	F	CN	F	mp.110°C

Ex. No.	R R R X	R ³	R ⁴	R ⁵	Physical properties
106	F ₂ CHCF ₂ CH ₃	F	CN	—о–снс <u>≕</u> сн	mp.в8°С
107	F ₃ C	F	CN	NH ₂	mp.193°C
108	F ₂ CHCF ₂ NCH ₃	F	CN	—о—сңс≘сн	mp.83°C
109	F ₃ C CH ₃	CI	CX	—о–снс <u></u> ен сн₃	mp 104°C
110	F ₃ C CH ₃	F	NO₂	F	mp.72°C
111	F ₃ C CH ₃	F	NO ₂	-0-ÇНС≣СН СН₃	mp.72°C

Ex. No.	R N N X	R ³	R ⁴	R ⁵	Physical properties
112	F ₃ C CH ₃	F	CN	-0СH ₂ С==СH ₂ И	mp.82°C
113	F ₃ C CH ₃	F	CN	-0-(H)	
114	F ₃ C CH ₃	F	CN	осн₂с <u>==</u> ссн₃	mp.138°C
115	F ₃ C	F	CN	-0-CH ₂ 0	mp. 72 ° C
116	F ₃ C CH ₃	F	CN	-0-(-) CH ₃	wax
117	F ₃ C CH ₃	F	CN	_0CH3	$n_{\rm D}^{20} = 1,5373$

ALCOHOLD VICE					
Ex. No.	R R R X	R ³	R ⁴	R ⁵	Physical properties
118	F ₃ C CH ₃	F	CN	-o-СH ₃	mp.121°C
119	F ₃ C CH ₃	F	CN	_o	mp.112°C
120	F ₃ C CH ₃	F	CN	-0	mp. 132°С
121	F ₃ C CH ₃	F	CN	-0-(mp.74°С
122	F ₃ C CH ₃	F	CN	_o	mp.45°C
123	F ₃ C CH ₃	F	CN	-0-()-F	mp.150°C

Ex. No.	R R R X	R³	R⁴	R ⁵	Physical properties
124	F ₃ C CH ₃	F	CN	-NHC₃H₁n	mp.124°C
125	F ₃ C CH ₃	F	СИ	-NHC ₂ H ₅	mp. 134°C
126	F ₃ C CH ₃	F	CN	NH ₂	mp.126°C
127	F ₂ CH CH ₃	F	CN	F	mp.116°C
128	F ₃ C CH ₃	F	CN	—о—сн ₂ —()	mp. 98°C
129	F ₃ C CH ₃	F	CN	-O-CH ₂ CH(CH ₃) ₂	mp.53°C

Ex. No.	R N N X	R ³	R ⁴	R ⁵	Physical properties
130	F ₃ C CH ₃	F	CN	O-C ₄ H ₉ n	mp.50°C
131	F ₃ C CH ₃	F	CN	-O-CH₂COOC₂H₅	mp. 214°C
132	F ₃ C CH ₃	F	CN	O-CHC₂H₅ CH₃	
133	F ₃ C CH ₃	F	CN	-0-CH ₂	mp.58°C
134	F ₃ C CH ₃	F	CN	\$\triangle \triangle \tria	mp.66°C
135	F ₃ C CH ₃	F	CN	—о—(н)	

Ex. No.	R R R X	R	R⁴	R ⁵	Physical Properties
136	F ₃ C CH ₃	F	СИ	OCH ₂ C=-CH ₃ CI	mp.53°C
137	F ₃ C CH(CH ₃) ₂	F	CN	F	n _D ²⁰ = 1,5012
138	F ₃ C NH ₂	F	CN	F	mp.69°С
139	F ₃ C CH ₃	F	CN	-O-CH ₂ CH=CH ₂	mp. 45°C
140	F ₃ C CH ₃	F	СИ	—осӊ₂с <u>==</u> сн	mp.99°C
141	F ₃ C CH ₃	F	CN	-OCH₂CH₂SC₂H₃	

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Ex. No.	R R X	R	R ⁴	R ^S	Physical properties
142	F ₃ C CH ₃	F	СИ	-O-CH ₂ Si(CH ₃) ₃	mp.89°C
143	F ₃ C CH ₃	F	CN	—о—снсңосн, сң	
144	F ₃ C CH(CH ₃) ₂	F	CN	-осн	mp.133°C
145	F ₃ C CH ₃	н	CN	СИ	mp.148°C
146	F ₃ C CH ₃	н	CN	CN	шр.78°С
147	F ₃ C NH ₂	Н	CN	F	mp.168°C

Ex. No.	R R X	R³	R ⁴	R ⁵	Physical properties
148	F ₃ C NH ₂	Н	CN	CN	mp.85°C
149	F ₃ C CH(CH ₃) ₂	Н	CN	СИ	mp.128°C
150	F ₃ C CH(CH ₃) ₂	Н	СИ	F	mp.76°C
151	F ₃ C CH(CH ₃) ₂	F	CN	—о—снс <u>≕</u> сн сн₃	
152	F ₃ C CH ₃	F	CN	OCH ₂ CF ₂ CF ₃	
153	F ₃ C , CH(CH ₃) ₂	F	CN	F	mp. 44°C

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Ex. No.	R R R X	R³	R ⁴	R ⁵	Physical properties
154	F ₃ C CH ₃	F	CN	_ocı	mp.111°C
155	F ₃ C CH ₃	Cl	CN	F	mp.110°C
156	F ₃ C CH ₃	F	č	-осн₄с <u>≕</u> ссн₃	mp.70°C
157	F ₃ C CH ₃	F	CN	-осн⁄сн=снсн₃	mp.57°C
158	F ₃ C CH ₃	F	CX	—осн ₂ —с <u>—</u> сн ₂ сн ₃	n _D ²⁰ = 1,5200
159	F ₃ C CH ₃	F	CN	—осн—сн=сн ₂ Сн ₃	n _D ²⁰ = 1,5149



Ex. No.	R R R X	R³	R⁴	R ⁵	Physical properties
160	F ₃ C CH ₃	F	CN	CH₃ -O-C-C≡CH CH₃	map. 89°C
161	F ₃ C CH ₃	F	CN	—осн ₂ сн ₂ с <u>—</u> сн ₂ сн ₃	mp. 80°C
162	F ₃ C CH ₃	F	CN	-OC₃H ₇ n	mp. 92°C
163	F ₃ C CH ₃	Cl	CN	—о—снс <u>≕</u> сн сн,	
164	F ₃ C NH ₂	н	CN	F ₃ C	mp.202°C
165	F ₃ C CH(CH ₃) ₂	Н	CN	F ₃ C , CH(CH ₃) ₂	mp.142°C

 $\widehat{\mathcal{G}}^{(k)}(x,y) \mapsto \sum_{i=1}^{k} \widehat{\mathcal{G}}_{i,i}(x,y) + \widehat{\mathcal{G}}_{i,i}(x,$

Ex. No.	R N X	R ³	R ⁴	R ⁵	Physical properties
166	F ₃ C CH ₃	F	CN	-0-()	mp.54°C
167	F ₃ C CH ₃	Н	CN	—оснс <u>≕</u> сн сн₃	mp. 140°C
168	F ₃ C CH ₃	F	č	-OCH(CH ₃)2	mp. 61°C
179	F ₃ C CH ₃	F	CN	—осн₂с <u>≕</u> сн	mp. 142°C
170	F ₃ C CH ₃	F	CN	оснсн _г осн _з / сн _з	
171	F ₃ C CH ₃	F	CN	-0	mp. 86°C

Section 1.					
Ex. No.	R R R X	R ³	R ⁴	R ⁵	Physical properties
172	F ₃ C CH ₃	F	CN	-OC₂H₅	mp.150°C
173	F ₃ C CH ₃	F	CN	-OC₄H₅n	mp.37°C
174	F ₃ C CH ₃	F	CN	Ş P	mp. 104°C
175	F ₃ C CH ₃	F	CZ	—OCHC₂H₅ CH₃	mp. 33°C
176	F ₃ C CH ₃	F	CN	-ОСН₂СН(СН₃)₂	mp. 79°C
177	F ₃ C CH ₃	F	CN	-OCH ₂ CH=CH ₂	mp.100°C

Ex. No.	R R R X	R ³	R ⁴	R ⁵	Physical properties
178	F ₃ C CH(CH ₃) ₂	Н	CN	—оснс <u>=</u> сн с́н₃	mp.108°C
179	F ₃ C C ₂ H ₅	F	CN	CI	mp. 53°C
180	F ₃ C C ₂ H ₅	F	СИ	—scн ₂ —	¹ H-NMR: 3,90-3,96; 4,20; 7,65-7,68.
181	F ₂ CHCF ₂ CH ₃	F	CN	F	mp. 85°C
182	F ₃ CCF ₂	F	CN	оснсңосн _з	¹ H-NMR: 1,38-1,40; 3,40; 4,57- 4,62, 7,40- 7,45.
183	F ₂ CH CH ₃	F	СИ	NH ₂	mp. 208°C

Ex. No.	R R R X	R³	R ⁴	R ⁵	Physical properties
184	F ₃ C CH ₃	F	CN	NH ₂	mp. 182°C
185	F ₃ C CH ₃	F	CN	S-CH ₂ -	mp. 77°C
186	F ₃ C CH ₃	F	CN	CI CH ₂ -C-CO ₂ CH ₃ CH ₃	oil
187	F ₃ C CH ₃	F	NO ₂	осн₂с≘сн	oil
188	CF ₃ C ₂ H ₅ N O	F .	CN	N(CH ₂ C≘CH) ₂	oil
189	CF ₃ C ₂ H ₅ N N O	F	CN	CH₂CCl₃	mp.114°C

Ex. No.	R R X	R³	!₹ ⁴	R ⁵	Physical properties
190	F ₃ C C ₂ H ₅	F	СИ	CI O CH ₂ -CH-C-NH✓	oil
191	F ₃ C C ₂ H ₅	F	CN	ОН	mp. 193°C
192	F ₃ C C ₂ H ₅	F	CN	CI CH ₂ -C-CO ₂ CH ₃ CH ₃	oil
193	F ₃ C C ₂ H ₅	F	CN	СІ СН₂-СН-СО₃СН₃	mp. 88°C
194	F ₃ C C ₂ H ₅	F	CN	ÇI CH₂−CH−CN	mp.140°C
195	F ₃ C C ₂ H ₅	F	CN	ÇI CH₂-CH-CO₂CH₃	oil

-111-

Ex. No.	R R X	R³	R ⁴	R ⁵	Physical properties
196	F ₂ HC CH ₃	F	CN	CI CH ₂ -C-CO ₂ CH ₃ CH ₃	mp. 113°C
197	F₂HC CH₃	F	CN	CI CH₂−CH−CO₂C₂H₅	oil

") The ¹H NMR spectra were recorded in deuterochloroform (CDCl₃) with tetramethylsilane (TMS) as the internal standard. The data given represent the chemical shift as 8 value in ppm.)

Use Examples:

5 In the use examples which follow, the compounds listed below were used as comparison substances:

$$H_3C$$
 CH_2 - $C\equiv CH$
 N
 O
 F
 CN
 (A)

3-Methyl-4-propargyl-1-(2,5-difluoro-4-cyano-phenyl)-1,2,4-triazolin-5-one

3,4-Dimethyl-1-(3-fluoro-4-cyano-phenyl)-1,2,4-triazolin-5-one (both disclosed in DE 3,839,480)

Example A

Pre-emergence test

Solvent:

5

10

15

5 parts by weight of acetone

Emulsifier:

1 part by weight of alkylaryl polyglycol

ether

To produce a suitable preparation of active compound, one part by weight of active compound is mixed with the stated amount of solvent, the stated amount of emulsifier is added and the concentrate is diluted with water to the desired concentration.

Seeds of the test plants are sown in normal soil and, after 24 hours, watered with the preparation of the active compound. It is expedient to keep constant the amount of water per unit area. The concentration of the active compound in the preparation is of no importance, only the amount of active compound applied per unit area being decisive. After three weeks, the degree of damage to the plants is rated in % damage in comparison to the development of the untreated control.

20 The figures denote:

0 % = no action (like untreated control)
100 % = total destruction

While Comparison Example (A) exhibits no herbicidal

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activity against weeds such as Setaria, Amaranthus, Chenopodium, Galinsoga, Matricaria, Solanum and Viola, at an application rate of 250 g/ha, activities between 40 and 100 % are shown, in this test, for example, by the compounds of Preparation Examples 7, 9, 17 and 29 and activities between 95 and 100 % by the compounds of Preparation Examples 10, 11, 12, 15 and 19.

Example B:

\$ 1 July 1 July 18 William (R) A Walter

5

15

20

Tetranychus test (OP resistant)

10 Solvent: 7 parts by weight of dimethylformamide
Emulsifier: 1 part by weight of alkylaryl polyglycol
ether

To produce a suitable preparation of active compound, l part by weight of active compound is mixed with the stated amount of solvent and the stated amount of emulsifier, and the concentrate is diluted with water to the desired concentrations.

Bean plants (Phaseolus vulgaris) which are severely infested with all developmental stages of the two-spotted spider mite (Tetranychus urticae) are dipped into a preparation of active compound at the desired concentration.

After the specified period of time, the mortality in per cent is determined . 100 %

means that all the spider mites have been killed; 0 \$ means that no spider mite has been killed.

In this test, a clearly superior acaricidal activity compared with Example (B), which is known from the prior art, is shown, for example, by compound 13 of the preparation examples.

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Example C

Phaedon-test

Solvent:

10

31 parts by weight of acetone

Emulsifier:

I part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, one part by weight of active compound is mixed with the stated amount of solvent, the stated amount of emulsifier is added and the concentrate is diluted with water to the desired concentration.

Cabbage leaves are treated with that suitable preparation of active compound. A such treated leave is put into a plastic box together with two Phaedon cochleariae in development stage. After 3 days an untreated leave is added. After the specified period of time, the mortality in per cent is determined. 100% means that all the Phaedon cochleariae have been killed; 0% means that no Phaedon cochleariae has been killed.

In this test a clearly superior acaricidal activity compared with the prior art is shown, for example, by compounds 20 and 62.

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Example D

Myzus-test

Solvent:

10

31 parts by weight of acetone

Emulsifier:

I part by weight of alkylaryl polyglycol ether

To produce a suitable preparation of active compound, one part by weight of active compound is mixed with the stated amount of solvent, the stated amount of emulsifier is added and the concentrate is diluted with water to the desired concentration.

Shoots of Vicia faba, which are stricken by Myzus persicae, are treated with such preparation of active compound in suitable concentration and put into a plastic box.

After the specified period of time the mortality in percent is determined. 100% means that all Myzus persicae have been killed; 0% means that no Myzus persicae has been killed.

15 In this test a clearly superior acaricidal activity in comparison to the prior art is shown for examles 57 and 62.

THE EMBODIMENTS OF THE INVENTION IN WHICH AN EXCLUSIVE PROPERTY OR PRIVILEGE IS CLAIMED ARE DEFINED AS FOLLOWS:

1. A substituted triazolinone of the general formula (I)

in which

R1 represents halogenoalkyl,

R² represents hydrogen, amino, cyano, alkyl, alkenyl, alkinyl, halogenoalkyl, halogenoalkenyl, halogenoalkinyl, alkoxyalkyl, alkylideneimino, or in each case optionally substituted cycloalkyl or cycloalkylalkyl,

R³ represents hydrogen or halogen,

R4 represents cyano or nitro,

 R^5 represents nitro, cyano, halogen, heterocyclylalkoxy, a radical of the formula R^6 , $-O-R^6$, $-S-R^6$, $-S(O)-R^6$, $-SO_2-R^6$, $-SO_2-O-R^6$, $-O-SO_2-R^6$, $-C(O)-O-R^6$, $-NR^6R^7$, $-SO_2-NR^6R^7$, $-C(O)-NR^6R^7$, $-NH-P(O)(OR^6)(R^7)$ or $-NH-P(O)(OR^6)(OR^7)$ or a radical of the formula

and

X represents oxygen or sulphur, where

 R^6 and R^7 independently of one another in each case represent hydrogen or in each case straight-chain or branched, optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl, cycloalkylalkyl, arylalkyl or aryl.

2. A substituted triazolinone of the general formula (I) according to claim 1, characterised in that

R¹ represents straight-chain or branched halogenoalkyl having 1 to 6 carbon atoms and 1 to 13 identical or different halogen atoms,

R² represents hydrogen, amino, cyano, straight-chain or branched alkyl having 1 to 8 carbon atoms, in each case straight-chain or branched alkenyl or alkinyl, each of which has 2 to 6 carbon atoms, straight-chain or branched halogenoalkyl having 1 to 6 carbon atoms and 1 to 13 identical or different halogen atoms, in each case straight-chain or branched halogenoalkenyl or halogenoalkinyl, each of which has 2 to 6 carbon atoms and 1 to 11 identical or different halogen atoms, straight-chain or branched alkoxyalkyl having 1 to 4 carbon atoms in each of the individual alkyl moieties, straight-chain or branched alkyl-



ideneimino having 1 to 8 carbon atoms, or cycloalkyl or cycloalkylalkyl, each of which has 3 to 8 carbon atoms in the cycloalkyl moiety and, if appropriate, 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, and each of which is optionally monosubstituted or polysubstituted in the cycloalkyl moiety by identical or different halogen substituents,

 $\ensuremath{\mathbb{R}}^3$ represents hydrogen, fluorine, chlorine, bromine or iodine,

R4 represents cyano or nitro,

 R^5 represents nitro, cyano, fluorine, chlorine, bromine, iodine or heterocyclyl- C_1 - C_4 -alkoxy, the heterocyclyl radical being represented by a three- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms, selected from oxygen and sulphur, or a radical of the formula R^6 , $-O-R^6$, $-S-R^6$, $-S(O)-R^6$ -, $-SO_2-R^6$, $-SO_2-O-R^6$, $-O-SO_2-R^6$, $-C(O)-O-R^6$, $-NR^6R^7$, $-SO_2-NR^6R^7$, $-C(O)-NR^6R^7$, $-NH-P(O)(OR^6)(R^7)$ or $-NH-P(O)(OR^6)(OR^7)$ or a radical of the formula

and

X represents oxygen or sulphur, where ${\tt R}^6$ and ${\tt R}^7$ independently of one another in each case represent hydrogen or straight-chain or branched alkyl which has

1 to 8 carbon atoms and which is optionally monosubstituted or polysubstituted by identical or different substituents, the substituents being:

halogen, in particular fluorine, chlorine, bromine and/or iodine, cyano, carboxyl, carbamoyl, in each case straight-chain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, alkoxycarbonylalkyl, N-alkyl-aminocarbonyl, cycloalkylaminocarbonyl, N,N-dialkylaminocarbonyl, trialkylsilyl or alkylsulphonylaminocarbonyl, each of which has 1 to 8 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl being represented by a five- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms, selected from nitrogen, oxygen and sulphur;

 ${\tt R}^6$ and ${\tt R}^7$ furthermore represent alkenyl or alkinyl, each of which has 2 to 8 carbon atoms and each of which is optionally monosubstituted or polysubstituted by identical or different halogen substitutents;

 R^6 and R^7 furthermore represent cycloalkyl which has 3 to 7 carbon atoms and which is optionally monosubstituted or polysubstituted by identical or different halogen substitutents, and/or by straight-chain or branched alkyl having 1 to 4 carbon atoms, or represent C_3 - C_7 -cycloalkyl- C_1 - C_3 -alkyl, or

R⁶ and R⁷ represent arylalkyl or aryl, each of which has 6 to 10 carbon atoms in the aryl moiety and 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety where present, and each of which is optionally monosubstituted or polysubstituted in the aryl moiety by identical or different

substituents, the aryl substituents in each case being:

halogen, cyano, nitro, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl, each of which has 1 to 6 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl, each of which has 1 to 6 carbon atoms and 1 to 13 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl, each of which has 1 to 6 carbon atoms in the individual alkyl moieties, or phenyl which is optionally monosubstitued or polysubstituted by identical or different halogen substituents and/or by straight-chain or branched alkyl or alkoxy, each of which has 1 to 6 carbon atoms, and/or by straight-chain or branched halogenoalkyl or halogenoalkoxy, each of which has 1 to 6 carbon atoms and 1 to 13 identical or different halogen atoms.

3. Substituted triazolinones of the general formula (I) according to claim 1, characterised in that

R¹ represents straight-chain or branched halogenoalkyl having 1 to 4 carbon atoms and 1 to 9 identical or different atoms of fluorine, chlorine or bromine,

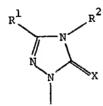
R² represents hydrogen, amino, cyano, straight-chain or branched alkyl having 1 to 6 carbon atoms, in each case straight-chain or branched alkenyl or alkinyl, each of which has 2 to 4 carbon atoms, straight-chain or branched halogenoalkyl having 1 to 4 carbon atoms and 1 to 9 identical or different atoms of fluorine, chlorine or bromine, in each case straight-chain or

branched halogenoalkenyl or halogenoalkinyl, each of which has 2 to 4 carbon atoms and 1 to 7 identical or different atoms of fluorine, chlorine or bromine, straight-chain or branched alkoxy-alkyl having 1 to 3 carbon atoms in each of the individual alkyl moieties, straight-chain or branched alkylideneimino having 1 to 6 carbon atoms, or cycloalkyl or cycloalkylalkyl, each of which has 3 to 7 carbon atoms in the cycloalkyl moiety and 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety, and each of which is optionally monosubstituted to tetrasubstituted in the cycloalkyl moiety by identical or different halogen substituents selected from fluorine, chlorine and bromine,

 ${\ensuremath{\mathsf{R}}}^3$ represents hydrogen, fluorine, chlorine or bromine,

R4 represents cyano or nitro,

 R^5 represents nitro, cyano, fluorine, chlorine, bromine or heterocyclyl- C_1 - C_3 -alkoxy, the heterocyclyl radical being represented by a four- or six-membered, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms selected from nitrogen, oxygen and sulphur, or a radical of the formula R^6 , $-O-R^6$, $-S-R^6$, $-S(O)-R^6$, $-SO_2-R^6$, $-SO_2-O-R^6$, $-O-SO_2-R^6$, $-C(O)-O-R^6$, $-NR^6R^7$, $-SO_2-NR^6R^7$, $-C(O)-NR^6R^7$, $-NH-P(O)(OR^6)(R^7)$ or $-NH-P(O)(OR^6)(OR^7)$ or a radical of the formula



and

X represents oxygen or sulphur, where

 ${\rm R}^6$ and ${\rm R}^7$ independently of one another in each case represent hydrogen or straight-chain or branched alkyl which has 1 to 6 carbon atoms and which is optionally monosubstituted, the substituents being:

cyano, carboxyl, carbamoyl, in each case straight-chain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, alkoxycarbonylalkyl, N-alkylaminocarbonyl, N,N-dialkylaminocarbonyl, trialkylsilyl or alkylsulphonylaminocarbonyl, each of which has 1 to 6 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl radical being represented by a five- or six-membered, saturated or unsaturated heterocycle having 1 to 3 identical or different hetero atoms selected from nitrogen, oxygen and sulphur;

 ${
m R}^6$ and ${
m R}^7$ furthermore represent straight-chain or branched halogenoalkyl having 1 to 4 carbon atoms and 1 to 9 identical or different atoms of fluorine, chlorine or bromine, and being optionally further substituted by ${
m C}_{1-2}$ -alkoxycarbonyl, ${
m C}_{1-6}$ -cycloalkylaminocarbonyl or cyano,

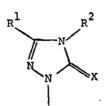
R⁶ and R⁷ furthermore represent alkenyl or alkinyl, each of which has 2 to 6 carbon atoms and each of which is optionally monosubstitued to trisubstituted by identical or different atoms of fluorine, chlorine or bromine;

 ${
m R}^6$ and ${
m R}^7$ furthermore represent cycloalkyl which has 3 to 6 carbon atoms and which is optionally monosubstituted to tetrasubstituted by identical or different atoms of fluorine, chlorine or bromine, and/or by straight-chain or branched alkyl having 1 to 3 carbon atoms, or represent ${
m C}_{3-6}$ -cycloalkyl- ${
m C}_{1-2}$ -alkyl, or represent phenylalkyl or phenyl, the first-mentioned

has 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety and each of which is optionally monosubstituted to trisubstituted in the phenyl moiety by identical or different substituents, the phenyl substituents in each case being:

halogen, cyano, nitro, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl, each of which has 1 to 4 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl, each of which has 1 to 4 carbon atoms and 1 to 9 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl, each of which has 1 to 4 carbon atoms in the individual alkyl moieties, or phenyl which is optionally monosubstituted or polysubstituted by identical or different halogen substituents and/or by straight-chain or branched alkyl or alkoxy, each of which has 1 to 4 carbon atoms, and/or by straight-chain or branched halogenoalkyl or halogenoalkoxy, each of which has 1 to 4 carbon atoms and 1 to 9 identical or different halogen atoms.

4. A compound according to claim 1, wherein



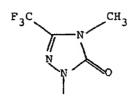
 R^3 is F, R^4 is CN and R^5 is H.

5. A compound according to claim 1, wherein

is

 ${\tt R}^3$ is F, ${\tt R}^4$ is CN and ${\tt R}^5$ is F.

6. A compound according to claim 1, wherein



 ${\mbox{R}}^3$ is F, ${\mbox{R}}^4$ is CN and ${\mbox{R}}^5$ is ${\mbox{CH}}_3$ -O-.

7. A compound according to claim 1, wherein

is

 R^3 is H, R^4 is CN and R^5 is F.

8. A compound according to claim 1, wherein

 R^3 is F, R^4 is CN and R^5 is CH_3 -O-.

9. A compound according to claim 1, wherein

is

 R^3 is F, R^4 is CN and R^5 is C_2H_5 -O-.

10. A compound according to claim 1, wherein

 R^3 is H, R^4 is CN and R^5 is F.

11. A compound according to claim 1, wherein

is

 ${\tt R}^3$ is F, ${\tt R}^4$ is CN and ${\tt R}^5$ is

12. A compound according to claim 1, wherein

$$R^3$$
 is F, R^4 is CN and R^5 is $--$ O-CH-C=CH CH₃

- 13. A pesticidal or herbicidal composition comprising a pesticidally or herbicidally effective amount of a compound according to any one of claims 1 to 12 in admixture with a suitable carrier or diluent.
- 14. A pesticidal or herbicidal composition comprising a pesticidally or herbicidally effective amount of a compound according to any one of claims 1 to 12 in admixture with a solid diluent or carrier, a liquified normally gaseous diluent or carrier, or a liquid diluent or carrier containing a surface active agent.
- 15. A method of combating pests or combating weeds which comprises applying to the pests or weeds, or to a habitat thereof, a pesticidally or herbicidally effective amount of a compound according to any one of claims 1 to 12.
- 16. A method of combating pests or combating weeds which comprises applying to the pests or weeds, or to a habitat

thereof, a pesticidally or herbicidally effective amount of a composition containing a compound according to any one of claims 1 to 12 in admixture with a suitable carrier or diluent.

- 17. A method of combating pests or combating weeds which comprises applying to the pests or weeds, or to a habitat thereof, a pesticidally or herbicidally effective amount of a composition containing between 0.0000001 and 95 % by weight of a compound according to any one of claims 1 to 12 in admixture with a suitable carrier or diluent.
- 18. A method of combating pests or combating weeds which comprises applying to the pests or weeds, or to a habitat thereof, a pesticidally or herbicidally effective amount of a composition containing between 0.0001 and 1 % by weight of a compound according to any one of claims 1 to 12 in admixture with a suitable carrier or diluent.
- 19. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a compound according to any one of claims 1 to 12 wherein the compound is applied as a pre-emergence herbicide.
- 20. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a compound according to any one of claims 1 to 12 wherein the compound is applied as a post-emergence herbicide.
- 21. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective

amount of a compound according to any one of claims 1 to 12 wherein the compound is applied to an area of cultivation at a rate of between 0.01 and 10 kg/ha.

- 22. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a compound according to any one of claims 1 to 12 wherein the compound is applied to an area of cultivation at a rate of between 0.05 and 5 kg/ha.
- 23. A process for preparing a compound of formula (I) as defined in claim 1, wherein R^1 , R^2 , R^3 , R^4 , R^5 and X are as defined in claim 1, which process comprises
 - a) reacting a 1H-triazolinone of the formula (II)

$$\begin{array}{c}
\mathbb{R}^{1} \\
\mathbb{N} \\
\mathbb{N}
\end{array}$$

$$\mathbb{R}^{2} \\
\mathbb{K}$$
(III)

in which

 ${\mbox{R}}^1, \ {\mbox{R}}^2$ and X have the above-mentioned meanings, with a halogenobenzene derivative of the formula (III)

$$\mathbb{R}^{3}$$

$$\mathbb{R}^{5}$$

$$\mathbb{R}^{5}$$

in which

 ${\tt R}^3$, ${\tt R}^4$ and ${\tt R}^5$ have the above-mentioned meanings and Hal represents halogen, or

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b) reacting a substituted triazolinone of the formula

(Ia)

$$\begin{array}{c}
\mathbb{R}^{1} \\
\mathbb{R}^{2} \\
\mathbb{R}^{3}
\end{array}$$

$$\mathbb{R}^{3} \\
\mathbb{R}^{5-1}$$
(Ia)

in which

 $\mbox{\ensuremath{\mbox{R}}}^1,\mbox{\ensuremath{\mbox{R}}}^2,\mbox{\ensuremath{\mbox{R}}}^4$ and X have the above-mentioned meanings and

 ${\tt R}^{5-1}$ represents halogen, with a nucleophile of the formula (IV)

$$R^{6-1}-Z-H \tag{IV}$$

in which

Z represents oxygen or sulphur and

 ${\tt R}^{6-1}$ represents in each case straight-chain or branched, optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl or aryl, and furthermore, in the event that Z represents oxygen, ${\tt R}^{6-1}$ also represents heterocyclyl, or

 $_{\mbox{\scriptsize c}})$ reacting a substituted triazolinone of the formula (Ib)

$$R^{1}$$
 R^{2-1}
 R^{3}
 R^{5}
(1b)

in which

 R^1 , R^3 , R^4 , R^5 and X have the above-mentioned meanings and

 $\ensuremath{\mathbb{R}^{2-1}}$ represents amino, with sodium nitrite in the presence of an acid or

 $\mbox{d)} \quad \mbox{reacting a substituted triazolinone of the} \\ \mbox{formula (Ic)} \\$

$$\begin{array}{c}
\mathbb{R}^{1} \\
\mathbb{R}^{2-2} \\
\mathbb{R}^{3}
\end{array}$$
(Ic)

in which

 ${\bf R}^1$, ${\bf R}^3$, ${\bf R}^4$, ${\bf R}^5$ and X have the above-mentioned meanings and

 ${\ensuremath{\mathbb{R}}}^{2-2}$ represents hydrogen, with an alkylating agent of the formula (V)

$$R^{2-3}-E \qquad (V)$$

in which

 $$\rm R^{2-3}$$ represents alkyl, alkenyl, alkinyl, halogenoalkyl, halogenoalkenyl, halogenoalkinyl, alkoxyalkyl or optionally substituted cycloalkyl and

E represents an electron-attracting leaving group.

- 24. A process for preparing a herbicidal or acaricidal composition comprising admixing a substituted triazolinone of the general formula (I) according to any one of claims 1 to 12 with an extender or surface-active agent.
- 25. 4-Amino-3-trifluoromethyl-lH-l,2,4-triazolin-5-one

26. 2,4-Difluoro-5-chlorobenzonitrile

27. A substituted triazolinone of the general formula (Ia)

$$\begin{array}{c}
\mathbb{R}^{1} \\
\mathbb{R}^{2} \\
\mathbb{R}^{3}
\end{array}$$

$$\mathbb{R}^{3} \\
\mathbb{R}^{5-1}$$

characterised in that

R¹ represents halogenoalkyl,

R² represents hydrogen, amino, cyano, alkyl, alkenyl, alkinyl, halogenoalkyl, halogenoalkenyl, halogenoalkinyl, alkoxyalkyl, alkylideneimino or in each case optionally substituted cycloalkyl or cycloalkylalkyl,

R³ represents hydrogen or halogen,

R4 represents cyano or nitro,

X represents oxygen or sulphur and

R⁵⁻¹ represents halogen.

28. A substituted triazolinone of the formula (Ib)



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characterised in that

R¹ represents halogenoalkyl,

 R^{2-1} represents amino,

R³ represents hydrogen or halogen,

R4 represents cyano or nitro,

 $\rm R^5$ represents nitro, cyano, halogen, heterocyclyloxy, a radical of the formula $\rm R^6$, $\rm -O-R^6$, $\rm -S-R^6$, $\rm -S(O)-R^6$, $\rm -SO_2-R^6$, $\rm -CO_2-O-R^6$, $\rm -CO_2-R^6$, $\rm -CO_3-R^6$, $\rm -CO_3-R^6$, $\rm -NR^6R^7$, $\rm -SO_2-NR^6R^7$, $\rm -CO_3-NR^6R^7$, $\rm -NH-P(O)(OR^6)(R^7)$ or $\rm -NH-P(O)(OR^6)(R^7)$ or a radical of the formula

and

X represents oxygen or sulphur, where

 ${\tt R}^6$ and ${\tt R}^7$ independently of one another in each case represent hydrogen or in each case straight-chain or branched, optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl or aryl.

29. A substituted triazolinone of the formula (Ic)





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(54) Substituted 1-Aryltriazolinones

(72) Linker, Karl-Heinz - Germany (Federal Republic of); Findeisen, Kurt - Germany (Federal Republic of); Haas, Wilhelm - Germany (Federal Republic of); Schallner, Otto - Germany (Federal Republic of); König, Klaus - Germany (Federal Republic of); Santel, Hans-Joachim - Germany (Federal Republic of); Dollinger, Markus - Germany (Federal Republic of);

- (71) Bayer Aktiengesellschaft Germany (Federal Republic of)
- (30) (DE) P 4309966.1 1993/03/26
- (57) 21 Claims

This application is as filed and may therefore contain an Notice: incomplete specification.

Industrie Canada Industry Canada

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Substituted 1-aryltriazolinones

Abstract

The invention relates to new substituted 1-aryltriazolinones of the general formula (I)

in which

R¹ represents hydrogen, alkyl, halogenoalkyl, alkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl or cycloalkyl,

R² represents a radical of the formula -NR³R³, R³ and R⁷ independently of one another in each case represent hydrogen, halogen, amino or nitro,

R⁵ represents nitro, cyano, halogen or halogenoalkyl, and

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- X represents oxygen or sulphur, where
- R^{8} represents hydrogen, alkyl, halogenoalkyl, a radical of the formula -CO- R^{12} or a radical of the formula -S(O)_n- R^{12} ,
- R^{5} represents alkyl, halogenoalkyl, a radical of the formula $-CO-R^{12}$ or a radical of the formula $-S(O)_{n}-R^{12}$,
- R¹⁰ represents hydrogen or represents in each case optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl, aryl, arylalkyl or heterocyclyl,
- R¹¹ represents hydrogen or represents in each case optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl, arylalkyl or aryl,
- R¹² represents in each case optionally substituted alkyl, cycloalkyl, arylalkyl, aryl or heterocyclyl, and
- n represents a number 0, 1 or 2,

to a number of processes for their preparation, to a number of new intermediates, and to their use as herbicides.

 J_i

The invention relates to new substituted 1-aryltriazolinones, to a number of processes for their preparation, to a number of new intermediates, and to their use as herbicides.

It is known that certain substituted triazolinones such as, for example, the compound 3-methyl-4-propargyl-1-(2,5-difluoro-4-cyano-phenyl)-1,2,4-triazolin-5-one possess herbicidal properties (cf. e.g. DE 38 39 480).

However the herbicidal activity of these previously known compounds with regard to problem weeds, and also their toleration by important crop plants, is not completely satisfactory in all areas of application.

New substituted 1-aryltriazolinones of the general formula (I),

15 have now been found in which

	R1	represents hydrogen, alkyl, halogenoalkyl,
		alkoxy, alkylthio, alkylsulphinyl, alkyl-
		sulphonyl or cycloalkyl,
	R ²	represents a radical of the formula -NR'R',
5	R³,	R ⁶ and R ⁷ independently of one another in each
		case represent hydrogen, halogen, amino or nitro
	\mathbb{R}^4	represents hydrogen, halogen, cyano or nitro,
		or one of the radicals $-R^{10}$, $-O-R^{10}$, $-S-R^{10}$,
		$-S(0)-R^{10}$, $-SO_2-R^{10}$, $-SO_2-OR^{10}$, $-SO_2-NR^{11}R^{10}$,
10		$-CO-OR^{10}$, $-CO-NR^{11}R^{10}$, $-O-SO_2-R^{10}$, $-N(R^{11})-SO_2-R^{10}$,
		$-NR^{11}R^{10}$, $-NH-P(O)(R^{11})(OR^{10})$ or
		$-NH-P(O)(OR^{11})(OR^{10}),$
	R ⁵	represents nitro, cyano, halogen or halogeno-
		alkyl, and
15	x	represents oxygen or sulphur, where
	R ⁴	represents hydrogen, alkyl, halogenoalkyl, a
		radical of the formula -CO-R12 or a radical of
		the formula $-S(O)_n-R^{12}$,
	R ⁹	represents alkyl, halogenoalkyl, a radical of
20		the formula -CO-R12 or a radical of the formula
		-S(0) _n -R ¹² ,
	R ¹⁰	represents hydrogen or represents in each case
		optionally substituted alkyl, alkenyl, alkinyl,
		cycloalkyl, aryl, arylalkyl or heterocyclyl,
25	R11	represents hydrogen or represents in each case
		optionally substituted alkyl, alkenyl, alkinyl,
		cycloalkyl, arylalkyl or aryl,
	R12	represents in each case optionally substituted
		alkyl, cycloalkyl, arylalkyl, aryl or hetero-
30		cyclyl, and

n represents a number 0, 1 or 2.

Depending on the nature of the substituents, the compounds of the formula (I) may possibly be present as geometrical and/or optical isomers or isomer mixtures of different composition. Both the pure isomers and the isomer mixtures are claimed according to the invention.

It has also been found that the new substituted 1-aryltriazolinones of the general formula (I),

in which

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- R¹ represents hydrogen, alkyl, halogenoalkyl, alkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl or cycloalkyl,
- R² represents a radical of the formula -NR⁹R⁹,
- R^3 , R^6 and R^7 independently of one another in each case represent hydrogen, halogen, amino or nitro,
- R^4 represents hydrogen, halogen, cyano or nitro, or one of the radicals $-R^{10}$, $-O-R^{10}$, $-S-R^{10}$, $-SO_2-R^{10}$, $-SO_2-OR^{10}$, $-SO_2-NR^{11}R^{10}$,

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		$-CO-OR^{10}$, $-CO-NR^{11}R^{10}$, $-O-SO_2-R^{10}$, $-N(R^{11})-SO_2-R^{10}$, $-NR^{11}R^{10}$, $-NH-P(O)(R^{11})(OR^{10})$ or $-NH-P(O)(OR^{11})(OR^{10})$,
	R ⁵	represents nitro, cyano, halogen or halogeno-
5		alkyl, and
	x	represents oxygen or sulphur, where
	R*	represents hydrogen, alkyl, halogenoalkyl, a
		radical of the formula -CO-R12 or a radical of
		the formula -S(O) _n -R ¹² ,
10	R9	represents alkyl, halogenoalkyl, a radical of
		the formula -CO-R12 or a radical of the formula
		$-S(0)_{n}-R^{12}$,
	R10	represents hydrogen or represents in each case
		optionally substituted alkyl, alkenyl, alkinyl,
15	-	cycloalkyl, aryl, arylalkyl or heterocyclyl,
	R11	represents hydrogen or represents in each case
		optionally substituted alkyl, alkenyl, alkinyl,
		cycloalkyl, arylalkyl or aryl,
	R12	represents in each case optionally substituted
20		alkyl, cycloalkyl, aryl, arylalkyl or hetero-
		cyclyl, and
	n	represents a number 0, 1 or 2

are obtained when

a) 1H-triazolinones of the formula (II),

in which

R1, R2 and X have the meaning given above,

are reacted with halogenobenzene derivatives of the formula (III),

5 in which

 $R^3\,,\ R^6\,,\ R^5\,,\ R^6$ and R^7 have the meanings given above and

Hal1 represents halogen,

optionally in the presence of a diluent and optionally in the presence of a reaction auxiliary, or when

b) substituted 1-aryltriazolinones of the formula (Ia),

in which

 $R^1,\ R^2,\ R^1,\ R^6,\ R^5,\ R^7$ and X have the meanings given above and

Hal' represents halogen,

5 are reacted with nucleophiles of the formula (IV),

in which

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 R^{13} represents a radical of the formula -0- R^{10} , -S- R^{10} or -NR¹¹ R^{10} , where R^{10} and R^{11} have the meanings given above,

optionally in the presence of a diluent and optionally in the presence of a reaction auxiliary, or when

c) substituted triazolinones of the formula (V),

in which

 \mathbb{R}^1 , \mathbb{R}^3 , \mathbb{R}^4 , \mathbb{R}^5 , \mathbb{R}^6 , \mathbb{R}^7 and X have the meanings given above

are reacted with alkylating, acylating or sulphonylating agents of the formula (VI),

$$R^9-E$$
 (VI)

in which

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- R' has the meaning given above and
- E represents an electron-attracting leaving group,

optionally in the presence of a diluent and optionally in the presence of a reaction auxiliary, or when

d) 4-alkylideneimino-triazolinones of the formula (VII),

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in which

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 R^1 , R^3 , R^4 , R^5 , R^6 , R^7 and X have the meanings given above,

 \mathbb{R}^{14} represents hydrogen or alkyl and \mathbb{R}^{15} represents alkyl or alkoxy,

are reacted with a reducing agent, optionally in the presence of a diluent and optionally in the presence of a reaction auxiliary.

Finally it has been found that the new substituted

10 1-aryltriazolinones of the general formula (I) possess
herbicidal properties.

Surprisingly, the substituted 1-aryltriazolinones of the general formula (I) according to the invention exhibit a considerably improved herbicidal activity against problem weeds with a comparable tolerance by crop plants in comparison to the substituted triazolinones known from

the state of the art, such as, for example, the compound 3-methyl-4-propargyl-1-(2,5-difluoro-4-cyanophenyl)-1,2,4-triazolin-5-one, which are closely related compounds in terms of their chemistry and their action.

- The general definition of the substituted 1-aryltriazolinones according to the invention is given by the formula (I). Preferred compounds of the formula (I) are those in which
- R1 represents hydrogen or represents in each case
 10 straight-chain or branched alkyl, alkoxy, alkylthio
 or alkylsulphonyl having in each case from 1 to 8
 carbon atoms, furthermore represents straight-chain
 or branched halogenoalkyl having from 1 to 8 carbon
 atoms and from 1 to 17 identical or different
 15 halogen atoms, or represents cycloalkyl having from
 3 to 8 carbon atoms,
 - R² represents a radical of the formula -NR⁶R⁹,
 - R³, R⁶ and R⁷ independently of one another in each case represent hydrogen, fluorine, chlorine, bromine, iodine, amino or nitro.
 - R⁴ represents hydrogen, fluorine, chlorine, bromine, iodine, cyano or nitro, or represents one of the radicals $-R^{10}$, $-O-R^{10}$, $-S-R^{10}$, $-S(O)-R^{10}$, $-SO_2-R^{10}$, $-SO_2-$
 - R⁵ represents nitro, cyano, fluorine, chlorine, bromine, iodine or represents straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and

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from 1 to 13 identical or different halogen atoms and

- X represents oxygen or sulphur, where
- represents hydrogen, straight-chain or branched alkyl having from 1 to 8 carbon atoms or straight-chain or branched halogenoalkyl having from 1 to 8 carbon atoms and from 1 to 17 identical or different halogen atoms, and furthermore represents a radical of the formula -CO-R¹² or a radical of the formula -S(O)_n-R¹²,
 - R° represents straight-chain or branched alkyl having from 1 to 8 carbon atoms or straight-chain or branched halogenoalkyl having from 1 to 8 carbon atoms and from 1 to 17 identical or different halogen atoms, and furthermore represents a radical of the formula -CO-R¹² or a radical of the formula -S(O)_n-R¹²,
 - R10 represents hydrogen;

- R10 furthermore represents straight-chain or branched
 20 <u>alkyl</u> having from 1 to 14 carbon atoms which is optionally substituted once or more than once by identical or different substituents, possible substituents being:
- halogen in particular fluorine, chlorine, bromine
 and/or iodine cyano, carboxyl, carbamoyl, in each
 case straight-chain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl,
 alkoxycarbonyl, N-alkylaminocarbonyl, N,N-dialkylaminocarbonyl or alkylsulphonylaminocarbonyl having
 in each case from 1 to 8 carbon atoms in the

individual alkyl moieties, or heterocyclyl, the heterocyclyl radical being a five- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur;

- R¹⁰ furthermore represents <u>alkenyl</u> or <u>alkinyl</u> having in each case from 2 to 8 carbon atoms, which are optionally substituted once or more than once by identical or different halogens in particular fluorine, chlorine, bromine and/or iodine;
- R10 represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once or more than once by identical or different substituents comprising halogen in particular fluorine, chlorine, bromine and/or iodine and/or straight-chain or branched alkyl having from 1 to 4 carbon atoms:
- R10 furthermore represents arylalkyl or aryl having in 20 each case from 6 to 10 carbon atoms in the aryl moiety and optionally from 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the aryl moiety once or more than once by identical or 25 different substituents, or represents a saturated or unsaturated, five- to seven-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur - which is optionally substituted once or once by identical 30 than different

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substituents and/or is benzo-fused, possible substituents of the aryl and/or heterocyclyl being in each case:

halogen, cyano, nitro, amino, N-acetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 6 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once or more than once by identical or different substituents comprising halogen and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 6 carbon atoms and/or straightchain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms;

R11 represents hydrogen;

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R¹¹ furthermore represents straight-chain or branched alkyl having from 1 to 14 carbon atoms which is optionally substituted once or more than once by identical or different substituents, possible substituents being:

halogen - in particular fluorine, chlorine, bromine and/or iodine - cyano, carboxyl, carbamoyl, in each

case straight-chain or branched alkoxy, alkoxy-alkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, N-alkylaminocarbonyl, N,N-dialkylaminocarbonyl or alkylsulphonylaminocarbonyl having in each case from 1 to 8 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl radical being a five- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur;

- R11 furthermore represents <u>alkenyl</u> or <u>alkinyl</u> having in each case from 2 to 8 carbon atoms, which are optionally substituted once or more than once by identical or different halogens in particular fluorine, chlorine, bromine and/or iodine;
- furthermore represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once or more than once by identical or different substituents comprising halogen in particular fluorine, chlorine, bromine and/or iodine and/or straight-chain or branched alkyl having from 1 to 4 carbon atoms;
- part furthermore represents arylalkyl or aryl having in each case from 6 to 10 carbon atoms in the aryl moiety and optionally 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the aryl moiety once or more than once by identical or different substituents, possible substituents of the

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aryl being in each case:

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R12

halogen, cyano, nitro, amino, N-acetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 6 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once or more than once by identical or different substituents comprising halogen and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 6 carbon atoms and/or straightchain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms; represents straight-chain or branched alkyl having from 1 to 8 carbon atoms which is optionally substituted once or more than once by identical or different substituents, possible substituents being: halogen - in particular fluorine, chlorine, bromine and/or iodine - cycloalkyl having from 3 to 8 carbon atoms or heterocyclyl, the heterocyclyl radical being a five- to seven-membered optionally benzofused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms - in

particular nitrogen, oxygen and/or sulphur;

R¹² furthermore represents <u>cycloalkyl</u> having from 3 to 7 carbon atoms which is optionally substituted once or more than once by identical or different substit-

uents comprising halogen - in particular fluorine, chlorine, bromine and/or iodine - and/or straight-chain or branched alkyl having from 1 to 4 carbon

atoms;

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furthermore represents <u>arylalkyl</u> or <u>aryl</u> having in each case from 6 to 10 carbon atoms in the aryl moiety and optionally from 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the aryl moiety once or more than once by identical or different substituents, or represents a saturated or unsaturated, five- to seven-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur - which is optionally substituted once or more than once by identical or different substituents, possible substituents of aryl or heterocyclyl being in each case:

halogen, cyano, nitro, amino, N-acetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 6 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or



different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once or more than once by identical or different substituents comprising halogen and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 6 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms and

n represents a number 0, 1 or 2.

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Particularly preferred compounds of the formula (I) are those in which

- R¹ represents hydrogen or in each case straight-chain or branched alkyl, alkoxy, alkylthic or alkylsulphonyl having in each case from 1 to 6 carbon atoms, or furthermore represents straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms in particular fluorine, chlorine or bromine or represents cycloalkyl having from 3 to 7 carbon atoms,
- 25 R² represents a radical of the formula -NR⁴R⁵,
 - R³, R⁶ and R⁷ independently of one another in each case represent hydrogen, fluorine, chlorine, bromine, amino or nitro,



R* represents hydrogen, fluorine, chlorine, bromine, cyano or nitro, or represents one of the radicals -R¹⁰, -O-R¹⁰, -S-R¹⁰, -S(O)-R¹⁰, -SO₂-R¹⁰, -SO₂-OR¹⁰, -SO₂-NR¹¹R¹⁰, -CO-OR¹⁰, -CO-NR¹¹R¹⁰, -O-SO₂-R¹⁰, -N(R¹¹)-SO₂-R¹⁰, -NR¹¹R¹⁰, -NH-P(O)(R¹¹)(OR¹⁰) or -NH-P(O)(OR¹¹)(OR¹⁰),

R⁵ represents nitro, cyano, fluorine, chlorine or bromine or represents straight-chain or branched halogenoalkyl having from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms - in particular fluorine, chlorine or bromine - and

X represents oxygen or sulphur, where

R⁶ represents hydrogen, straight-chain or branched alkyl having from 1 to 6 carbon atoms or straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms - in particular fluorine, chlorine or bromine - and furthermore represents a radical of the formula -CO-R¹² or a radical of the formula -S(O)_n-R¹²,

R' represents straight-chain or branched alkyl having from 1 to 6 carbon atoms or straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms - in particular fluorine, chlorine or bromine - and furthermore represents a radical of the formula -CO-R¹² or a radical of the formula -S(O)_n-R¹²,

R10 represents hydrogen;

30 R¹⁰ furthermore represents straight-chain or branched

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alkyl having from 1 to 12 carbon atoms which is optionally substituted once or twice by identical or different substituents, possible substituents being: cyano, carboxyl, carbamoyl, in each case straightchain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, N-N, N-dialkylaminocarbonyl alkylaminocarbonyl, alkylsulphonylaminocarbonyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl radical being a five- to seven-membered, optionally benzofused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur; furthermore represents straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms in particular fluorine, chlorine and/or bromine, furthermore represents alkenyl or alkinyl having in each case from 2 to 6 carbon atoms, which are in each case optionally substituted once to three times by identical or different halogens - in particular fluorine, chlorine and/or bromine; furthermore represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once to three times by identical or different substituents comprising halogen - in particular fluorine, chlorine and/or bromine - and/or straight-chain or

30 R¹⁰ furthermore represents phenylalkyl or phenyl having

branched alkyl having from 1 to 3 carbon atoms;

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R10

R10

R10

optionarly from 1 to 3 carbon atoms in the straightchain or branched alkyl moiety, which are in each case optionally substituted in the phenyl moiety once to five times by identical or different substituents, or represents a saturated or unsaturated, five- to seven-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur - which is optionally substituted once to three times by identical or different substituents and/or is benzofused, possible substituents of phenyl or heterocyclyl being in each case: fluorine, chlorine, bromine, cyano, nitro, amino, Nacetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 4 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 4 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once to five times by identical or different substituents comprising fluorine, chlorine, bromine and/or straightchain or branched alkyl or alkoxy having in each case from 1 to 4 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having

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in each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms;

R11 represents hydrogen;

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R11 furthermore represents straight-chain or branched alkyl having from 1 to 12 carbon atoms which is optionally substituted once or twice by identical or different substituents, possible substituents being: cyano, carboxyl, carbamoyl, in each case straightchain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, Nalkylaminocarbonyl, N, N-dialkylaminocarbonyl alkylsulphonylaminocarbonyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl radical being a five- to seven-membered, optionally benzofused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur;

R¹¹ furthermore represents straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms in particular fluorine, chlorine and/or bromine,

R¹¹ furthermore represents <u>alkenyl</u> or <u>alkinyl</u> having in each case from 2 to 6 carbon atoms, which are in each case optionally substituted once to three times by identical or different halogens - in particular fluorine, chlorine and/or bromine;

R¹¹ furthermore represents <u>cycloalkyl</u> having from 3 to 7 carbon atoms which is optionally substituted once to three times by identical or different

substituents comprising halogen - in particular fluorine, chlorine and/or bromine - and/or straight-chain or branched alkyl having from 1 to 3 carbon atoms;

furthermore represents phenylalkyl or phenyl having optionally from 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the phenyl moiety once to five times by identical or different substituents, possible substituents of phenyl being in each case:

fluorine, chlorine, bromine, cyano, nitro, amino, Nacetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 4 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 4 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once to five times by identical or different substituents comprising fluorine, chlorine, bromine and/or straightchain or branched alkyl or alkoxy having in each case from 1 to 4 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 4 carbon atoms and from 1 to

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		9 identical or different halogen atoms;
	R12	represents straight-chain or branched alkyl having
		from 1 to 12 carbon atoms which is optionally
		substituted once or twice by identical or different
5		substituents, possible substituents being:
		cycloalkyl having from 3 to 7 carbon atoms or
		heterocyclyl, the heterocyclyl radical being a five-
		to seven-membered, optionally benzo-fused, saturated
		or unsaturated heterocycle having from 1 to 3
10		identical or different hetero atoms - in particular
		nitrogen, oxygen and/or sulphur;
	R12	furthermore represents halogenoalkyl having from 1
		to 6 carbon atoms and from 1 to 13 identical or
		different halogen atoms - in particular fluorine,
15		chlorine and/or bromine,
	R12	furthermore represents cycloalkyl having from 3 to
		7 carbon atoms which is optionally substituted once
		to three times by identical or different substit-

uents comprising halogen - in particular fluorine, chlorine and/or bromine - and/or straight-chain or branched alkyl having from 1 to 3 carbon atoms;

R¹² furthermore represents <u>phenylalkyl</u> or <u>phenyl</u> having optionally from 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the phenyl moiety

once to five times by identical or different substituents, or represents a saturated or unsaturated, five- to seven-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms -in particular nitrogen, oxygen and/or sulphur - which

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is optionally substituted once to three times by identical or different substituents and/or is benzo-fused, possible substituents of phenyl or heterocyclyl being in each case:

fluorine, chlorine, bromine, cyano, nitro, amino, Nacetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 4 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 4 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once to five times by identical or different substituents comprising fluorine, chlorine, bromine and/or straightchain or branched alkyl or alkoxy having in each case from 1 to 4 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms and

25 n represents a number 0, 1 or 2.

Very particularly preferred compounds of the formula (I) are those in which

R1 represents hydrogen or represents in each case

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straight-chain or branched alkyl, alkoxy, alkylthio or alkylsulphonyl having in each case from 1 to 4 carbon atoms, or furthermore represents straight-chain or branched halogenoalkyl having from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms - in particular fluorine, chlorine or bromine - or represents cycloalkyl having 3, 5 or 6 carbon atoms.

- R2 represents a radical of the formula -NR4R9,
- 10 R³, R⁶ and R⁷ independently of one another in each case represent hydrogen, fluorine, chlorine, bromine, amino or nitro,

 - R⁵ represents nitro, cyano, fluorine, chlorine or bromine, or straight-chain or branched halogenoalkyl having from 1 to 3 carbon atoms and from 1 to 7 identical or different halogen atoms, and
 - X represents oxygen or sulphur, where
- R⁶ represents hydrogen, straight-chain or branched
 25 alkyl having from 1 to 4 carbon atoms or straightchain or branched halogenoalkyl having from 1 to 4
 carbon atoms and from 1 to 9 identical or different
 halogen atoms in particular fluorine, chlorine or
 bromine and furthermore represents a radical of
 the formula -CO-R¹² or a radical of the formula

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-S(0),-R12,

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R' represents straight-chain or branched alkyl having from 1 to 4 carbon atoms or straight-chain or branched halogenoalkyl having from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms - in particular fluorine, chlorine or bromine - or furthermore represents a radical of the formula -CO-R¹² or a radical of the formula -S(O)_n-R¹²,

R10 represents hydrogen;

10 R¹⁰ furthermore represents straight-chain or branched alkyl having from 1 to 8 carbon atoms which is optionally substituted once, possible substituents being:

cyano, carboxyl, carbamoyl, in each case straightchain or branched alkoxy, alkoxyalkoxy, alkylthio,
alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, Nalkylaminocarbonyl, N,N-dialkylaminocarbonyl or
alkylsulphonylaminocarbonyl having in each case from
1 to 4 carbon atoms in the individual alkyl
moieties, or heterocyclyl, the heterocyclyl radical
being a five- or six-membered, optionally benzofused, saturated or aromatic heterocycle having from
1 to 3 identical or different hetero atoms - in
particular nitrogen, oxygen and/or sulphur;

25 R¹⁰ furthermore represents straight-chain or branched <u>halogenoalkyl</u> having from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms -in particular fluorine, chlorine and/or bromine,

R¹⁰ furthermore represents <u>alkenyl</u> or <u>alkinyl</u> having in each case from 2 to 6 carbon atoms, which are in



each case optionally substituted once to three times by identical or different halogens - in particular fluorine, chlorine and/or bromine;

furthermore represents <u>cycloalkyl</u> having 3, 5 or 6 carbon atoms which is optionally substituted once to three times by identical or different substituents comprising halogen - in particular fluorine, chlorine and/or bromine - and/or straight-chain or branched alkyl having from 1 to 3 carbon atoms;

furthermore represents phenylalkyl or phenyl having optionally from 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the phenyl moiety once to three times by identical or different substituents, or represents a saturated or aromatic, five- or six-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur - which is optionally substituted once to three times by identical or different substituents and/or is benzofused, possible substituents of phenyl or heterocyclyl being in each case:

fluorine, chlorine, bromine, cyano, nitro, amino, Nacetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 3 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, haogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 3 carbon atoms and

R10

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from 1 to 7 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 3 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once to three times by identical or different substituents comprising fluorine, chlorine, bromine and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 3 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case 1 to 3 carbon atoms and 1 to 7 identical or different halogen atoms;

R11 represents hydrogen;

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furthermore represents straight-chain or branched alkyl having from 1 to 8 carbon atoms which is optionally substituted once, possible substituents being:

cyano, carboxyl, carbamoyl, in each case straightchain or branched alkoxy, alkoxyalkoxy, alkylthio,
alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, Nalkylaminocarbonyl, N,N-dialkylaminocarbonyl or
alkylsulphonylaminocarbonyl having in each case from
1 to 4 carbon atoms in the individual alkyl
moieties, or heterocyclyl, the heterocyclyl radical
being a five- or six-membered, optionally benzofused, saturated or aromatic heterocycle having from
1 to 3 identical or different hetero atoms - in
particular nitrogen, oxygen and/or sulphur;

R¹¹ furthermore represents straight-chain or branched halogenoalkyl having from 1 to 4 carbon atoms and



from 1 to 9 identical or different halogen atoms -in particular fluorine, chlorine and/or bromine,

- R¹¹ furthermore represents <u>alkenyl</u> or <u>alkinyl</u> having in each case from 2 to 6 carbon atoms, which are in each case optionally substituted once to three times by identical or different halogens - in particular fluorine, chlorine and/or bromine;
- R¹¹ furthermore represents <u>cycloalkyl</u> having 3, 5 or 6 carbon atoms which is optionally substituted once to three times by identical or different substituents comprising halogen in particular fluorine, chlorine and/or bromine and/or straight-chain or branched alkyl having from 1 to 3 carbon atoms;
- R¹¹ furthermore represents <u>phenylalkyl</u> or <u>phenyl</u> having optionally from 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the phenyl moiety once to three times by identical or different substituents, possible substituents of phenyl being in each case:

fluorine, chlorine, bromine, cyano, nitro, amino, Nacetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 3 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 3 carbon atoms and from 1 to 7 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl

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or alkoximinoalkyl having in each case from 1 to 3 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once to three times by identical or different substituents comprising fluorine, chlorine, bromine and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 3 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case 1 to 3 carbon atoms and 1 to 7 identical or different halogen atoms;

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R¹² represents straight-chain or branched <u>alkyl</u> having from 1 to 4 carbon atoms which is optionally substituted once, possible substituents being: cycloalkyl having 3, 5 or 6 carbon atoms or heterocyclyl, the heterocyclyl radical being a five- or six-membered, optionally benzo-fused, saturated or aromatic heterocycle having from 1 to 3 identical or

different hetero atoms - in particular nitrogen,

oxygen and/or sulphur;

20 R¹² furthermore represents straight-chain or branched halogenoalkyl having from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms - in particular fluorine, chlorine and/or bromine,

furthermore represents cycloalkyl having 3, 5 or 6 carbon atoms which is optionally substituted once to three times by identical or different substituents comprising halogen - in particular fluorine, chlorine and/or bromine - and/or straight-chain or branched alkyl having from 1 to 3 carbon atoms;

30 R12 furthermore represents phenylalkyl or phenyl having



optionally from 1 to 3 carbon atoms in the straightchain or branched alkyl moiety, which are in each case optionally substituted in the phenyl moiety once to three times by identical or different substituents, or represents a saturated or aromatic, five- or six-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms - in particular nitrogen, oxygen and/or sulphur - which is optionally substituted once to three times by identical or different substituents and/or is benzofused, possible substituents of phenyl or heterocyclyl being in each case: fluorine, chlorine, bromine, cyano, nitro, amino, Nacetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 3 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 3 carbon atoms and from 1 to 7 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 3 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once to three times by identical or different substituents comprising fluorine, chlorine, bromine and/or straightchain or branched alkyl or alkoxy having in each

case from 1 to 3 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having

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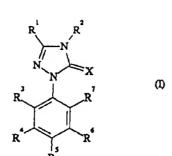


in each case from 1 to 3 carbon atoms and from 1 to 7 identical or different halogen atoms and represents a number 0, 1 or 2.

Individually, and apart from the compounds listed in the Preparation Examples, the following substituted triazolinones of the general formula (I) may be mentioned:

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R ¹	R²	R³	R ⁴	R ⁵	R ⁶	R7	x
CH ₃	-NH-CH ₃	H	ОН	CN	Н	F	0
CH ₃	-NH-CH ₃	H	OH	CN	Н	а	0
CH ₃	-NH-CH ₃	H	OH	NO ₂	H	F	0
CH ₃	-NH-CH ₃	H	OH	NO ₂	H	а	0
CH ₃	-NH-CH ₃	H	OH	CF ₃	H	a	0
CH ₃	-NH-CH ₃	а	OH	NO_2	H	а	0
CH_3	-NH-CH ₃	NO_2	OH	CF ₃	H	NO_2	O
CH_3	-NH-CH ₃	H	-O-CH3	CN	H	F	0
CH_3	-NH-CH ₃	H	-O-CH ₃	CF3	α	H	0
CH ₃	-NH-CH ₃	H	-O-CH3	NO_2	H	F	0

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\mathbb{R}^1	R²	R³	R ⁴	R ⁵	R ⁶	R7	x	
CH ₃	-NH-CH ₃	Н	a	CN	Н	Н	0	_
CH ₃	-NH-CH ₃	Н	-O-CH(CH ₃)-COOC ₂ H ₅	CN	H	a a	_	
CH ₃	-NH-CH ₃	H	-O-CH ₂ -CN	CN	H	F	0	
CH ₃	-NH-CH ₃	Н	_	CN	H	D.	0	
.,			-0	CIN	п	u	0	
CU-	NIII OII							
CH ₃	-NH-CH ₃	H	-0-0-0	NO_2	H	F	Ο.	
CH ₃	-NH-CH ₃	H	-O-CH2-C(CI)=CH2	CN	Н	H	0	
CH ₃	-NH-CH ₃	H	-O-CH2-C(CI)=CH2	CN	н	а	0	
CH ₃	-NH-CH ₃	F	F	CF ₃	H	H	0	
CH ₃	-NH-CH ₃	H	-O-SO ₂ -CH ₃	CN	H .	F	0	
CH ₃	-NH-CH ₃	H	-O-SO ₂ -CH ₃	CN	Н	a	0	
CH ₃	-NH-CH ₃	H	-O-SO ₂ -CH ₃	NO_2	Н	a	0	
				_				
CH_3	-NH-CH ₃	H	F	CN	H	H	0	
CH ₃	-NH-CH ₃	H	-O-CHF ₂	CN	H	F	0	
CH3	-NH-CH3	Н	-O-CHF ₂	CN	H	α	0	
CH ₃	-NH-CH ₃	H	-O-CHF ₂	NO ₂	H	F	0	
CH ₃	-NH-CH ₃	H	-O-CHF ₂	CF3	H	F	0	
CH ₃	-NH-CH ₃	H	-O-CHF ₂	NO_2	H	α	0	
СН3	-NH-CH ₃	H	-SCH ₃	NO ₂	H	α	0	
CH ₃	-NH-CH ₃	H	-SCH ₃	NO ₂	н	F	0	



R1	R²	R³	R ⁴	R ⁵	R'	R7	x	
CH ₃	-NH-CH ₃	н	-SCH3	CN	Н	F	0	
CH ₃	-NH-CH3	H	-SCH ₃	CN	Н	H	0	
CH ₃	-NH-CH3	H	-SCH ₃	CF ₃	H	а	· 0	
CH3	-NH-CH ₃	а	-SCH ₃	CF ₃	Н	а	0	
CH ₃	-NH-CH ₃	H	-NH-CH ₃	CN	H	а	0	
CH ₃	-NH-CH ₃	H	-NH-CH ₃	CN	H	F	0	
CH ₃	-NH-CH ₃	H	-NH-CH ₃	CN	H	H	О	
CH ₃	-NH-CH ₃	H	-NH-CH ₃	NO_2	H	F	0	
CH ₃	.,	a	-NH-CH ₃	CF ₃	H	a	0	
CH ₃	-NH-CH ₃	a	-N(CH ₃) ₂	CF ₃	н	a	0	
CH ₃	-NH-CH ₃	H	-N(CH ₃) ₂	CN	H	a	0	
CH ₃	-NH-CH ₃	H	-N(CH ₃) ₂	CN	H	F	0	
CH ₃	-NH-CH ₃	Н	-N(CH ₃) ₂	NO_2	H	a	0	
						_	_	
CH ₃	-NH-CH3	H	-NH-CH ₂ -CH=CH ₂	NO_2	H .	F	0	
CH ₃	-NH-CH ₃	H	-NH-CH ₂ -CH=CH ₂	CN	H	a	0	
CH ₃	-NH-CH ₃	H	-NH-SO ₂ -CH ₃	NO ₂	H	a	0	
CH ₃	-NH-CH3	H	-N(CH ₃)-SO ₂ -CH ₃	NO ₂	H	a	0	
CH_3	-NH-CH3	H	-COOC ₂ H ₅	CN	H	F	0	
CH_3	-NH-CH ₃	H	-COOCH ₃	CN	H	α	0	
CH ₃	-NH-CH ₃	H	-CO-NH-CH ₃	CN	H	α	O	
CH_3	-NH-CH ₃	H	-CO-N(CH ₃) ₂	CN	H	F	0	
CH ₃	-NH-CH ₃	H	-S(O)-CH3	CN	H	F	0	

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R1	R²	R ³	R ⁴	R ⁵	R ⁶	R ⁷	x
CH ₃	-NH-CH ₃	н	-SO ₂ -CH ₃	NO ₂	Н	F	0
CH ₃	-NH-CH ₃	H	-SO ₂ -NH-CH ₃	NO ₂	Н	а	0
CH ₃	-NH-CH ₃	H	-SO ₂ -NH-CH ₃	CN	Н	F	0
CH ₃	-NH-CH ₃	H	-SO ₂ -O-CH ₃	CN	н	F	0
СН3	-NH-CH ₃	Н	O NHPOCH, CH,	СŃ	Н	F	O
СН3	-NH-CH3	Н	O -NH-P(OC ₂ H ₂) ₂	CN	н	a	0
C ₂ H ₅	-NH-CH ₃	а	H	CN	Н	CI	0
C ₂ H ₅	-NH-CH ₃	NO ₂	н	CF ₃	H	NO_2	0
C ₂ H ₅	-NH-CH ₃	Н	СН3	NO ₂	H	H	0
C_2H_5	-NH-CH ₃	H	C ₂ H ₅	NO ₂	H	H	0
C_2H_5	-NH-CH ₃	a	, F	CF ₃	H	а	0
	-NH-CH3	H	а	CN	а	H	0
	-NH-CH ₃		-OCH ₃	CN	H	F	0
	-NH-C ₂ H ₅		-OCH ₃	CN	H	F	0
	-NH-C ₂ H ₅		F	CF3	H	a	0
	-NH-C ₂ H ₅		-NH-SO ₂ -CH ₃	CN	H	F	0
	-NH-C ₂ H ₅		-O-CH(CH ₃)-C≡CH	CN	H	F	0
	-NH-C ₂ H ₅		-S-C ₂ H ₅	CN	H	F	0
n-C ₃ H	7 -NH-CH	3 NC) ₂ H	CF ₃	H	NO_2	0



R1	R³	R³	R ⁴	R ⁵	R ⁴	R'	х	
n-C ₃ H ₇	-NH-CH	3 H	ОН	CN	H	a	0	
n-C ₃ H ₇	-NH-CH	, H	-OCH3	CN	H	F	0	
n-C ₃ H ₇	-NH-CH	, н	-NH-SO2-CH3	CN	Ħ	а	0	
n-C3H7	-NH-CH	H	-O-CH(CH3)-C≡CH	CN	н	F	0	
n-C3H7	-NH-CH3	з Н	-S-CH(CH ₃)-COOCH ₃	CN	Н	F	0	
n-C3H7	-NH-CH3	Н	-O-CH ₂ -COOCH ₃	CN	H	F	0	
n Calla	-NH-CH ₂	11	CIV.	NO	77			
., ,	• •		CH ₃	NO ₂	H	H	0	
i-C ₃ H ₇	-NH-CH ₃		СН ₃	NO ₂	H	H	0	
i-C3H7	-NH-CH ₃	H	, · · · · · · · · · · · · · · · · · · ·	CN	H	F	0	
			-o- ⟨ ⟩-a					
i-C3H7	-NH-CH ₃	H	H	NO ₂	H	H	0	
i-C3H7	-NH-CH ₃	H	-COOC ₂ H ₅	CN	H	F	0	
i-C ₃ H ₇	-NH-CH ₃	H	-CO-NH-CH3	CN	H	а	0	
i-C ₃ H ₇	-NH-CH ₃	Н	-S-C ₂ H ₅	CN	H	F	0	
i-C3H7	-NH-CH ₃	Н	-S-(CH ₂) ₂ -OC ₂ H ₅	CN	Н	F	0	
	-NH-CH3	NO ₂	H	CF ₃	H	NO ₂	0	
> − ·	-NH-CH3	H	ОН	CN	H.	F	0	
<u></u>	NH-CH-	ш	ഹസ്ദ	CN	ч	F	0	



R1	R ²	R³	R ⁴	R ⁵	R ⁶	R ⁷	x	
>	-NH-CH3	н	-CO-NH-CH ₃	CN	Н	F	0	
<u></u>	-NH-CH ₃	Н	-O-SO ₂ -CH ₃	CN	Н	F	0	
\triangleright	-NH-CH ₃	Н	-80 ₂ -О-СН ₃	CN	H	F	0	
>	-NH-CH ₃	н	-NH-SO ₂	CN	н	F	o	
	-NH-СН3		-o-ar, o	CN	н	F	0	
<u></u>	-NH-CH ₃	Н	-0-CH ₂ S	CN	н	F	o	
>	-NH-CH ₃	н —с	o-Cai-coc	L CN	н	F	0	
<u></u>	-N(CH2)2	NO2	— сн, н	CEo		NO.	0	



R ¹	R ² R	.	R ⁴	R5	R*	R ⁷	x
<u>×</u>	-N(CH ₃) ₂		н	CF3	Н	Н	0
\triangleright	-N(CH ₃) ₂	Н	ОН	CN	н	СІ	0
<u></u>	-NH-CH3	н	-OCH ₃	CN	н	F	0
\triangleright	-NH-CH3	н	-S-C ₂ H ₅	CN	н	F	0
<u></u>	-NH-CH ₃	Н	-N(CH ₃) ₂	CN	н	F	0
	-NH-CH ₃	Н	-O-CH(CH ₃)-COO-C ₂ H ₅	CN	H	F	0
>	-NH-CH ₃	н	-SO ₂ -OCH ₃	CN	Н	F	
>	-N(CH ₃) ₂	н	о —nh-р(ос _г н _{э)} ,	CN	н	F	0
СН3	-N(CH ₃) ₂	H	0 NH-POCH, CH,	CN	Н	F	0



R1	R ²	R³	R ⁴	R ⁵	R ⁶	R7	x
СН3	-N(CH ₃) ₂	Н	ОН	CN	Н	F	0
CH ₃	-N(CH ₃) ₂	H	ОН	CN	Н	а	0
CH ₃	-N(CH ₃) ₂	H	OH	NO_2	H	a	0
CH ₃	$-N(CH_3)_2$	H	ОН	NO_2	H	F	0
CH ₃	-N(CH ₃) ₂	H	-NH ₂	CN	H	F	0
CH ₃	-N(CH ₃) ₂	H	-NH ₂	CN	H	a	0
CH ₃	$-N(CH_3)_2$	H	-OCH ₃	CN	H	F	0
CH ₃	-N(CH ₃) ₂	H	-OCH ₃	CN	н	F	S
	.v.or.		a av	on r	••	_	
CH ₃	-N(CH ₃) ₂	H	-S-CH ₃	CN	H	F	0
CH ₃	$-N(CH_3)_2$	Н	-S-CH ₃	CN	H	F	S
CH ₃	$-N(CH_3)_2$	H	-CO-NH-CH ₃	CN	H	F	0
CH_3	-N(CH3)2	H	-CO-N(CH ₃) ₂	CN	H	а	0
CH ₃	$-N(CH_3)_2$	H	$-NH-SO_2-CH_3$	CN	H.	a	S
CH ₃	$-N(CH_3)_2$	H	-NH-SO2-C2H5	CN	H	F	S
CH ₃	-N(CH3)2	H	-OCHF ₂	CN	H	F	0
CH_3	-N(CH3)2	H	-OCHF ₂	CN	H	F	0
CH ₃	$-N(CH_3)_2$	H	$-N(CH_3)_2$	CN	H	F	0
CH ₃	-N(CH ₃) ₂	H	-NH-C ₂ H ₅	CN	H	F	0
CH ₃	$-N(CH_3)_2$	H	$-O-CH_2-C(Cl)=CH_2$	CN	H	F	0
CH ₃	$-N(CH_3)_2$	H	-O-SO ₂ -CH ₃	CN	H	F	0
CH ₃	$-N(CH_3)_2$	H	-SO ₂ -O-CH ₃	CN	H	а	0



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R1	R²	R³	R ⁴	R ⁵	R ⁶	R7	x
СН3	-N(CH ₃) ₂	Н	-0-	CN	Н	F	0
CH_3	-N(CH ₃) ₂	H	-O-CH ₂ -C ₆ H ₅	CN	H	F	0
СН3	-N(CH ₃) ₂	H	-0- - F	CN	H	F	0
CH ₃	-N(CH ₃) ₂	Н	-0-{F	CN	Н	F	0
СН3	-N(CH ₃) ₂	Ħ	-O-CH(CH ₃)-COOCH ₃	CN	н	F	0
CH ₃	-N(CH ₃) ₂	H	-O-CH(CH ₃)-COOCH ₃	NO ₂	н	F	0
CH3	-N(CH ₃) ₂	а	-O-CH(CH ₃)-COOCH ₃	CF3	Н	a	0
CH ₃	-N(CH ₃) ₂	a	-O-CH(CH3)-C≡CH	CF ₃	H	a	0
CH ₃	-N(CH ₃) ₂	H	-O-CH(CH3)-C≣CH	NO ₂	H	F	0
C_2H_5	$-N(CH_3)_2$	H	-O-CH(CH3)-C≡CH	NO_2	H	F	0
C ₂ H ₅	-N(CH ₃) ₂	Н	-O-CH ₂ -C≡CH	NO ₂	H	F	0
Calla	NI/CTI-N-	Ť.T	0.60- 6-0-	CN	ч	F	0
C ₂ H ₅	-N(CH ₃) ₂	H H	-O-CH ₂ -C ₆ H ₅		H	F	0
C ₂ H ₅	-N(CH ₃) ₂		-N(CH ₃) ₂	CN		_	_
C ₂ H ₅	-N(CH ₃) ₂	H	-N(CH ₃) ₂	NO ₂	H	F	0
C ₂ H ₅	-N(CH ₃) ₂	H	-OCH ₃	NO ₂	H	F	0
C ₂ H ₅	-N(CH ₃) ₂	H	-OCH ₃	CN	H	a	0
C ₂ H ₅	-N(CH ₃) ₂	H	OH	CN	H	a	0
CH_3	-N(CH ₃) ₂	H	SH	CN.	H	F	0





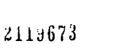
R1	R ²	R³	R ⁴	R ⁵	R ⁵	R ⁵ R ⁷	
CH ₃	-N(CH ₃) ₂	Н	SH	CN	Н	F	0
i-C3H7	-N(CH ₃) ₂	H	SH	CN	H	F	0
i-C3H7	-N(CH ₃) ₂	H	OH	CN	H	F	0
i-C3H7	-N(CH ₃) ₂	H	OCH ₃	CN	H	F	0
_i C ₃ H ₇	-N(CH ₃) ₂	NO ₂	H	CF3	H	NO_2	0
i-C3H7	$-N(CH_3)_2$	а	H	CN	H	a	0
i-C ₃ H ₇	-N(CH ₃) ₂	H	CI	CN	CI	H	0
i-C3H7	-N(CH ₃) ₂	H	-S-CH(CH ₃)-COOCH ₃	CN	H	F	0
i-C3H7	$-N(CH_3)_2$	H	-O-CH ₂ -CN	CN	H	F	0
i-C3H7	-N(CH3)2	H	-O-C(CI)=CH ₂	CN	H	F	0
i-C3H7	-N(CH3)2	H	-COOCH3	CN	H	F	0
i-C3H7	$-N(CH_3)_2$	H	$-N(CH_3)-SO_2-CH_3$	CN	H	F	0
i-C ₃ H ₇	$-N(CH_3)_2$	H	$-N(CH_3)-SO_2-C_2H_5$	CN	Н	F	0
i-C ₃ H ₇	-N(CH3)2	H	-CO-NH-CH ₃	CN	H	F	0
i-C3H7	$-N(CH_3)_2$	H	$-N(C_2H_5)-SO_2-CH_3$	CN	H	а	0
	$-N(CH_3)_2$	H	$-NH-SO_2-CH_3$	CN	H	F	0
	$N(CH_3)_2$		-O-CH(CH3)-C≡CH	CN	H	F	0
	N(CH ₃) ₂	H	-O-CH ₂ -CH=CH ₂	CN	H	F	0
	$N(CH_3)_2$	H	-0-CH ₂ -C≡CH	CN	H	F	0
	$N(CH_3)_2$	H	$-NH-SO_2-C_2H_5$	CN	H	F	0
CH ₃	N(CH ₃) ₂	a	-NH-SO ₂ -CH ₃	NO_2	H	a	0

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2	

R1	R ²	R³	R ⁴	R ⁵	R ⁴	R7	х
CH ₃	-N(CH ₃) ₂	H	-NH ₂	CN	Н	F	0
CH ₃	-N(CH ₃) ₂	H	OH	CN	H	F	0
CH3	-N(CH ₃) ₂	H	H	CN	Ĥ	F	0
CH ₃	-N(CH ₃) ₂	H	SH	CN	H	F	Ο
CH ₃	-N(CH ₃) ₂	H	-OCH3	CN	H	F	0
CIT.	NAME OF THE PERSON OF THE PERS						
CH ₃	-N(CH ₃) ₂	H	-O-CH(CH ₃)-C≡CH	CN	H	F	0
CH ₃	-N(CH ₃) ₂	Н	-O-CH ₂ -CH=CH ₂	CN	H	F	0
CH ₃	-N(CH ₃) ₂	H	-O-CH ₂ -C≡CH	CN	H	a	0
CH ₃	-N(CH ₃) ₂	H	-NH-SO ₂ -C ₂ H ₅	CN	H	F	0
CH ₃	-N(CH ₃) ₂	H	-NH-SO ₂ -CH ₃	CN	н	F	0
CF ₃	-NH-CH ₃	H	F	CN	H	а	S
CF ₃	-NH-CH ₃	H	OH	CN	H	F	0
CF_3	-NH-CH ₃	H	OH	NO ₂	H	F	0
CF ₃	-NH-CH ₃	H	-OCH3	CN	H	F	0
CF3	-NH-CH ₃	H	-OCH ₃	NO_2	H	F	0
CF3	-NH-CH ₃	H	-OCH ₃	CN	H	F	0
CF3	-NH-CH ₃	H	-O-CH ₂ -CN	CN	H	F	0
CF ₃	-NH-CH ₃	H	-COOCH ₃	CN	H	F	0
CF ₃	-NH-CH ₃	H	-COOCH ₃	CN	H	а	0
CF ₃	-NH-CH ₃	H	-N(CH ₃)-SO ₂ -C ₂ H ₅	CN	H	а	0



R1	R²	R³	R ⁴	R ⁵	R ⁶	R7	x
CF ₃	-NH-CH ₃	NO_2	Н	CF ₃	Н	NO ₂	0
CF ₃	-NH-CH3	H	a	CN	а	н	0
CF ₃	-NH-CH ₃	H	-O-CH ₂ -CH=CH ₂	CN	н	F	0
CF_3	-NH-CH ₃	H	-NH-CH ₂ -CH=CH ₂	CN	H	F	0
F ₂ CH-	-NH-CH ₃	Н	F	CN	7.7	<i>~</i>	
F ₂ CH-	_		OH		H	a	0
F ₂ CH-	••		-O-CH ₂ -CH=CH ₂	CN	H	F	0
F ₂ CH-	.,			CN	H	F	0
12011-	-Nn-Cng	н	-O-CH ₂ -CH=CH ₂	CN	H	F	S
F ₂ CH-	-NH-CH ₃	H	-NH-SO ₂ -n-C ₄ H ₉	CN	H	F	0
F ₂ CH-	-NH-CH ₃	H	-S-C ₂ H ₅	CN	H	F	0
F ₂ CH-	-N(CH ₃) ₂	H	-S-C ₂ H ₅	CN	H	F	0
F ₂ CH-	-N(CH ₃) ₂	H	$-NH-SO_2-C_2H_5$	CN	H	а	0
F ₂ CH-	-N(CH ₃) ₂	H	CH ₃	NO_2	H	H	0
F ₂ CH-	-N(CH ₃) ₂	H	-O-CH ₂ -C≡CH	NO_2	H	F	0
F ₂ CH-	-N(CH ₃) ₂		-COOC ₂ H ₅	CN	H	F	0
F ₂ CH-	-N(CH ₃) ₂		-S-CH(CH ₃)-COOCH ₃	CN	H	F	0
CF ₃	-N(CH ₃)		-NH-SO ₂ -CH ₃	NO_2	H	α	0
CF ₃	-N(CH ₃)		-O-SO ₂ -CH ₃	CN	H	F	0
CF ₃	-N(CH ₃)		-SO ₂ -CH ₃	CN	Ħ	F	0

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R1	R ² R ²	1	R ⁴	R ⁵	R ⁴	R'	х
CF ₃	-N(CH ₃) ₂	H	-NH-CH(CH ₃) ₂	CN	H	а	0
CF ₃	-N(CH ₃) ₂	H	-O-CH2-C≡CH	CN	H	F	0
CF_3	-N(CH ₃) ₂	H	-COOC ₂ H ₅	CN	Н	F	0
CF ₃	$-N(CH_3)_2$	H	-CO-NH-CH ₃	CN	H	F	0
CF3	$-N(CH_3)_2$	H	-O-SO ₂ -CH ₃	CN	H	а	0
CH ₃ -S-	-N(CH ₃) ₂	н	-CH ₃	NO ₂	Н	H	0
	-N(CH ₃) ₂	н	ОН	CN	Н	F	0
-	-N(CH ₃) ₂	H	ОН	CN	Н	а	0
CH3-S-	-N(CH ₃) ₂	Н	-S-C ₂ H ₅	CN	н	F	o
CH3-S-	-N(CH ₃) ₂	H	-O-CH ₂ -C≡CH	NO ₂	H	F	0
CH3-S-	$-N(CH_3)_2$	H	-COOCH ₃	CN	H	F	0
CH3-S-	$-N(CH_3)_2$	H	$-NH-CH(CH_3)_2$	CN	H	F	0
CH3-\$-	$-N(CH_3)_2$	H	-N(CH ₃)-SO ₂ -CH ₃	CN	H	F	0
CH3-S-	-NH-CH3	H	-S-C ₂ H ₅	CN	H	F	0
CH ₃ -S-	-NH-CH ₃	H	-NH-CH2-CH=CH2	CN	H	F	0
CH3-\$-	-NH-CH ₃	H	-O-CH ₂ -CN	CN	H	F	o
CH ₃ -S-	-NH-CH ₃	H	-O-SO ₂ -CH ₃	CN	H	F	0

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R1	R²	R ³		R ⁴ R	5	R ⁶	R ⁷	x	
CH ₃	-S-	-NH-CH ₃	н	-O-CH(CH ₃)-СООСН	3 CN	Н	F	0	
CH ₃	·s-	-NH-CH ₃	H	-O-СН ₂ -С _б Н ₅	CN	H	F	0	
CH ₃	·\$-	-NH-CH ₃	H	-O-CH2-C6H5	NO ₂	Н	F	0	
CH ₃ -	·S-	-NH-CH3	H	-COOCH3	CN	Н	CI	0	
СН3-	·S-	-NH-CH3	H		CN	H	F	0	
				-0-CH2 0					
CH3-S(O)-	-N(CH ₃) ₂	Н	-CH3	NO ₂	н	н	0	
CH3-S	(O)-	-N(CH ₃) ₂	Н	ОН	CN	H	F	0	
CH3-S(O)-	-N(CH ₃) ₂	H	OH	CN	H	а	0	
CH3-S(O)-	-N(CH ₃) ₂	Н	-S-C ₂ H ₅	CN	H	F	0	
CH3-S(O)-	-N(CH ₃) ₂	H	-O-CH ₂ -C≡CH	NO ₂	H	F	0	
CH ₃ -S(O)-	-N(CH ₃) ₂	H	-COOСН3	CN	H	F	0	
CH3-S(0)-	-N(CH ₃) ₂	H	-NH-CH(CH ₃) ₂	CN	H	F	0	
CH3-S(0)-	-N(CH ₃) ₂	H	-N(CH ₃)-SO ₂ -CH ₃	CN	H	F	0	
CH3-S(0)-	-NH-CH ₃	H	-S-C ₂ H ₅	CN	H	F	0	
CH3-S(0)-	-NH-CH ₃	Н	-NH-CH ₂ -CH=CH ₂	CN	Н	F	0	
CH3-S(O)-	-NH-CH ₃	H	-O-CH ₂ -CN	CN	H	F	0	
CH3-S(0	O)-	-NH-CH ₃	H	-O-SO ₂ -CH ₃	CN	H	. F	0	
CH3-S(())-	-NH-CH ₃	H	-O-CH(CH ₃)-COOCH ₃	CN	Н	F	0	
CH3-S(0)-	-NH-CH ₃	H	-O-CH ₂ -C ₆ H ₅	CN	H	F	0	
CH3-S(0))-	-NH-CH ₃	H	-O-CH ₂ -C ₆ H ₅	NO ₂	H	F	0	
CH3-S(0))-	-NH-CH ₃	H	-COOCH3	CN	H	a	0	





R ¹ R	2 R3		R ⁴ R	5	R ⁶	R ⁷	x	
CH3-S(O)	NH-CH3	H	-0-CI-1 0	CN	H	F	0	
CH3-SO2-	-N(CH ₃) ₂	Н	-CH ₃	NO ₂	Н	н		
CH3-SO2-	-N(CH ₃) ₂	H	_	CN	Н	F	0	
CH ₃ -SO ₂ -	-N(CH ₃) ₂	H	ОН	CN	н	CI	0	
CH3-SO2-	-N(CH ₃) ₂	H	-S-C ₂ H ₅	CN	H	F	0	
CH ₃ -SO ₂ -	-N(CH ₃) ₂	H	-O-CH ₂ -C≘CH	NO_2	H	F	0	
CH ₃ -SO ₂ -	-N(CH ₃) ₂	H	-COOCH3	CN	H	F	0	
CH3-SO2-	-N(CH ₃) ₂	Н	-NH-CH(CH ₃) ₂	CN	H	F	0	
CH3-SO2-	-N(CH ₃) ₂	H	-N(CH ₃)-SO ₂ -CH ₃	CN	H	F	0	
CH ₃ -SO ₂ -	-NH-CH ₃	H	-S-C ₂ H ₅	CN	Н	F	0	
CH ₃ -SO ₂ -	-NH-CH ₃	Н	-NH-CH2-CH=CH2	CN	H	F	O	
CH ₃ -SO ₂ -	-NH-CH ₃	H	-O-CH ₂ -CN	CN	H	F	0	
CH3-SO2-	-NH-CH ₃	H	-O-SO ₂ -CH ₃	CN	H	F	0	
CH3-SO2-	-NH-CH ₃	H	-O-CH(CH ₃)-COOCH ₃	CN	H	F	0	
CH_3-SO_2-	-NH-CH ₃	H	-O-CH ₂ -C ₆ H ₅	CN	H	F	0	
CH ₃ -SO ₂ -	-NH-CH ₃	H	-O-CH ₂ -C ₆ H ₅	NO_2	H	F	0	
CH ₃ -SO ₂ -	-NH-CH ₃	H	-COOCH3	CN	H	a	0	
CH3-SO2-	-NH-CH ₃	H		CN	H	F	0	
			-0-CH2 0					

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R1	R ²	R³	R4	R ⁴ R ⁵		R7	х
CH ₃ O	-N(CH		2 H	CF ₃	Н	NO ₂	0
CH ₃ O	-N(CH		F	CN	H	H	0
CH ₃ O	-N(CH		a	CN	a	Н	0
CH ₃ O	-N(CH ₂		-OCH ₃	CN	H	F	0
CH ₃ O	-N(CH ₃) ₂ H	-S-C ₂ H ₅	CN	H	F	0
CH ₃ O	-N(CH ₃)2 H	-O-CH ₂ -CH=CH ₂	CN	H	a	0
- Сн ₃ 0	-N(CH ₃)2 H	NW.			_	
CH ₃ O	-N(CH ₃		NH ₂	CN	H	F	0
CH ₃ O			-N(CH ₃)-SO ₂ -CH ₃	CN	H	F	0
CH ₃ O	-N(CH ₃)		-COOCH3	CN	H	F	0
-	-N(CH ₃)		H	NO_2	H	a	0
CH ₃ O	-N(CH ₃)		-S-CH(CH ₃)-COOCH	3 CN	H	a	0
CH ₃ O	-N(CH ₃)		-O-CH(CI)=CH ₂	CN	H	F	0
CH ₃ O	-NH-CH	3 H	OH	CN	H	F	0
CH30	-NH-CH	3 H	OH	NO_2	H	F	0
CH ₃ O	-NH-CH	3 H	-OCH3	NO_2	H	F	0
CH ₃ O	-NH-CH	3 H	-OCH ₃	CN	H	a	0
CH ₃ O	-NH-CH	3 H	-SCH ₃	CN	H	a	0
CH ₃ O	-NH-CH	3 H	-O-CH(CH3)-C≡CH	CN	H	F	0
CH ₃ O	-NH-CH	3 H	-O-CH(CH3)-C≡CH	CN	H	F	S
CH ₃ O	-NH-CH	3 H	-O-CH2-CH=CH2	CN	H	F	0

R1	R²	R³		R ⁴	R ⁵	R'	R ⁷	x
CH ₃ O	-NH-	СН3	Н	-NH-CH2-CH	I=CH ₂ CN	Н	F	0
CH ₃ O	-NH-	CH_3	H	-NH-CH2-CH	I=CH ₂ NO ₂	H	F	0
CH ₃ O	-NH-	CH_3	H	-NH-SO ₂ -C	2H ₅ NO ₂	H	F	0
CH ₃ O	-NH-	CH3	H	-NH-SO ₂ -C	₂ H ₅ CN	H	а	0
CH_3O	-NH-	CH3	H	-NH-SO ₂ -C	2H5 CN	H	F	0
CH ₃ O	-NH-	CH ₃	H	-NH-SO ₂ -C	₂ H ₅ CN	H	F	S
CH ₃	-NH-	CF3	H	F	CN	H	H	0
CH_3	-NH-	CF3	H	-OCH ₃	CN	Н	F	0
CH_3	-NH-	CF3	H	-S-C ₂ H ₅	CN	H	F	0
CH ₃ CH ₃ CH ₃ CH ₃ CH ₃ CH ₃	-NH-(-NH-(-NH-(-NH-(-NH-(F ₃ F ₃ F ₃ F ₃	н н н н	-O-CH(CH ₃)-(-NH-SO ₂ -C ₂ -NH-SO ₂ -C ₂ -NH-SO ₂ -C ₂ -NH-SO ₂ -C ₂ -COOCH ₂	H ₅ CN H ₅ CN H ₅ CN H ₅ CN	H H H H	F F Cl F	0 0 0 0 0
СН ₃ СН ₃ Н Н	-NH-C -NH-C -N(CH	F ₃ 3)2 3)2	H H H	-NH-CH ₃ OH F OH	CZ CZ CZ CZ	H H H	F F F	0 0 0
H	-N(CH	3)2	H	-O-CH(CH ₃)-C	ECH CN	H	F	0
H	-N(CH		H	-NH-SO ₂ -CH		H	F	o



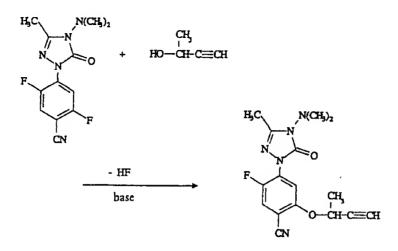
R	1 R ² F	٤,	R ⁴ F	5	R [¢]	R ⁷	x	
н	-N(CH ₃) ₂	H	-N(CH ₃) ₂	CN	Н	F	0	
H	-N(CH ₃) ₂	H	-N(CH ₃)SO ₂ -C ₂ H ₅	CN	H	F	0	
H	-N(CH ₃) ₂	H	-NH-SO ₂ -C ₂ H ₅	CN	H	а	0	
H	-N(CH ₃) ₂	a	F	CF ₃	H	а	0	
H	-NH-CH ₃	H	-O-CH(CH3)-C≡CH	CN	H	F	0	
H	-NH-CH ₃	H	-NH-SO ₂ -C ₂ H ₅	CN	H	F	0	
H	-NH-CH ₃	H	-O-CH(CH ₃) ₂	CN	H	F	0	
H	-NH-CH ₃	H	-O-CH(CH ₃)-CH ₂ -OCH ₃	CN	Н	F	0	
H	-NH-CH ₃	H	-S-C ₂ H ₅	CN	H	F	0	
H	-NH-CH3	H	-O-(CH ₂) ₂ -NH-CH ₃	CN	H	F	0	
H	-NH-CH3	H	OH	CN	Ή	F	0	
H	-NH-CH3	H	-O-CH ₃	CN	H	F	0	
H	-NH-CH3	H	SH	CN	H	F	0	
H	-NH-CH ₃	H	-S-C ₂ H ₅	CN	Н	F	0	
H	-NH-CH ₃	H	-NH-SO ₂ -CH ₃	CN	H	F	0	
H	-NH-CH ₃	H	-O-CH ₂ -C <u>≡</u> CH	CN	H	F	0	
H	-N(CH ₃) ₂	H	OH	CN	H	F	0	
H	$-N(CH_3)_2$	H	-O-C ₂ H ₅	CN	H	а	0	
H	-NH-CH3	H	-S-C ₂ H ₅	CN	H	a	0	
H	-N(CH ₃) ₂	H	-NH-SO ₂ -CH ₃	CN	H	α	0	
H	-N(CH ₃) ₂	H	~	CN	Н	F	0	
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Using, for example, 3-methyl-4-dimethylamino-1,2,4-triazolin-5-one and 2,4,5-trifluorobenzonitrile as starting materials, the sequence of reaction of process (a) according to the invention can be represented by the following formula scheme:

Using, for example, 1-(4-cyano-2,5-difluorophenyl)-3-methyl-4-dimethylamino-1,2,4-triazolin-5-one and 1-butin-3-ol as starting materials, the sequence of reaction of process (b) according to the invention can be represented by the following formula scheme:

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Using, for example, 1-(4-cyano-2-fluoro-5-methoxy-phenyl)-4-amino-3-methyl-1,2,4-triazolin-5-one as the starting compound and acetyl chloride as the acylating agent, the sequence of reaction of process (c) according to the invention can be represented by the following formula scheme:



Using, for example, 1-(4-cyano-2-fluoro-5-methoxy-phenyl)-3-methyl-4-isopropylideneimino-1,2,4-triazolin-5-one as the starting compound and sodium borohydride as the reducing agent, the sequence of reaction of process (d) according to the invention can be represented by the following formula scheme:

A general definition of the lH-triazolinones required as starting materials for carrying out process (a) according to the invention is given by the formula (II). In this formula (II), R¹, R² and X preferably and particularly preferably represent those radicals which have already been mentioned as preferred and particularly preferred for these substituents in connection with the description of the compounds of the formula (I) according to the invention.

The 1H-triazolinones of the formula (II) are known or are obtainable by analogy with known processes (cf. e.g. Chimica Acta Turcica 9, 381 [1981]; EP 399 294; EP 422 469; J. Heterocycl. Chem. 10, 387-390 [1973]; Indian J. Chem. 7, 959-963 [1969]; DE 37 19 575; DE 38 03 523; Liebigs Ann. Chem. 637, 135 [1960]; J. Heterocycl. Chem.

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16, 403 [1979]; J. Heterocycl. Chem. 17, 1691 [1980]; J. Indian Chem. Soc. 57, 270-272 [1980]; Indian J. Chem. Sect. B 22B, 270-271 [1983]; Chem. Ber. 98, 3025 [1965]; JP 52-125168; Europ. J. Med. Chem. 18, 215 [1983]).

A general definition of the halogenobenzene derivatives furthermore required as starting materials for carrying out process (a) according to the invention is given by the formula (III). In this formula (III), R³, R⁴, R⁵, R⁶ and R⁷ preferably and particularly preferably represent those radicals which have already been mentioned as preferred and particularly preferred for these substituents in connection with the description of the compounds of the formula (I) according to the invention. Hal¹ represents preferably fluorine, chlorine or bromine, in particular fluorine or chlorine.

The halogenobenzene derivatives of the formula (III) are generally known or are obtainable by analogy with known processes (cf. e.g. EP 191 181; EP 441 004; EP 431 373). The compound 5-chloro-2,4-difluorobenzonitrile is not already known. It is obtained by reacting the known compound 2,4,5-trichlorobenzonitrile (cf. e.g. EP 441 004) with potassium fluoride, optionally in the presence of a diluent such as, for example, tetramethylene sulphone at temperatures of between 100°C and 200°C (compare also in this respect the Preparation Examples).

A general definition of the substituted triazolinones required as starting materials for carrying out process (b) according to the invention is given by the formula (Ia). In this formula (Ia), R¹, R², R³, R⁴, R⁵, R⁷ and X preferably and particularly preferably represent those radicals which have already been mentioned as being preferred and particularly preferred for these substituents in connection with the description of the substances of the formula (I) according to the invention. Hal² represents preferably fluorine, chlorine or bromine, in

The substituted triazolinones of the formula (Ia) compounds according to the invention and obtainable by means of processes (a), (c) and/or (d) according to the invention.

particular fluorine or chlorine.

A general definition of the nucleophiles furthermore required as starting materials for carrying out process (b) according to the invention is given by the formula (IV). In this formula (IV), R¹³ preferably represents a radical of the formula -O-R¹⁰, -S-R¹⁰ or -NR¹¹R¹⁰, where R¹⁰ and R¹¹ preferably represent those radicals which have already been mentioned as being preferred and particularly preferred for these substituents in connection with the description of the substances of the formula (I) according to the invention. The nucleophiles of the formula (IV) are generally known compounds of organic chemistry.

A general definition of the substituted triazolinones required as starting materials for carrying out process (c) according to the invention is given by the formula (V). In this formula (V), R¹, R³, R⁴, R⁵, R⁶, R⁷ and X

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preferably and particularly preferably represent those radicals which have already been mentioned as being preferred and particularly preferred for these substituents in connection with the description of the substances of the formula (I) according to the invention.

Substituted triazolinones of the formula (V) are not already known. They are, however, to a large extent the subject of the Applicant's as yet unpublished patent applications and are obtainable by means of the processes described therein, for example by reacting 4-amino-litriazolinones of the formula (VIII),

in which

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R1 has the meaning given above,

with halogenobenzene derivatives of the formula (III),

15 in which



 \mathbb{R}^3 , \mathbb{R}^4 , \mathbb{R}^5 , \mathbb{R}^6 and \mathbb{R}^7 have the meaning given above and Hal¹ represents halogen,

optionally in the presence of a diluent and optionally in the presence of a reaction auxiliary, in analogy to the implementation of process (a) according to the invention.

4-Amino-1H-triazolinones of the formula (VIII) are known or are obtainable by analogy with known processes (cf. e.g. EP 294 666; J. Heterocycl. Chem. 10, 387-390 [1973]; Indian J. Chem. 7, 959-963 [1969]; DE 37 19 575; DE 38 03 523; Liebigs Ann. Chem. 637, 135 [1960]; J. Heterocycl. Chem. 16, 403 [1979]; J. Heterocycl. Chem. 17, 1691 [1980]; J. Indian Chem. Soc. 57, 270-272 [1980]; Indian J. Chem. Sect. B 22B, 270-271 [1983]; Chem. Ber. 98, 3025 [1965]; JP 52-125168; Europ. J. Med. Chem. 18, 215 [1983]).

A general definition of the alkylating, acylating and sulphonylating agents furthermore required as starting materials for carrying out process (c) according to the invention is given by the formula (VI). In this formula (VI), R' represents preferably and particularly preferably those radicals which have already been mentioned as preferred and particularly preferred for these substituents in connection with the description of the substances of the formula (I) according to the invention. E represents a conventional electron-attracting leaving radical such as, for example, halogen, in particular chlorine, bromine or iodine or, in the case of the alkylating

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agents, represents in each case optionally substituted alkylsulphonyloxy, alkoxysulphonyloxy or arylsulphonyloxy, oxy, such as, in particular, methanesulphonyloxy, trifluoromethanesulphonyloxy, methoxysulphonyloxy, ethoxysulphonyloxy or p-toluenesulphonyloxy.

The alkylating, acylating and sulphonylating agents of

The alkylating, acylating and sulphonylating agents of the formula (VI) are generally known compounds of organic chemistry.

A general definition of the substituted 4-alkylideneimino-triazolinones required as starting materials for carrying out process (d) according to the invention is given by the formula (VII). In this formula (VII), R¹, R³, R⁴, R⁵, R⁶, R⁷ and X preferably and particularly preferably represent those radicals which have already been mentioned as being preferred and particularly preferred for these substituents in connection with the description of the substances of the formula (I) according to the invention. R¹⁴ represents preferably hydrogen or straightchain or branched alkyl having from 1 to 4 carbon atoms, in particular hydrogen, methyl or ethyl. R¹⁵ represents preferably in each case straight-chain or branched alkyl or alkoxy having in each case from 1 to 4 carbon atoms, in particular methyl, ethyl, methoxy or ethoxy.

The 4-alkylideneimino-triazolinones of the formula (VII) are not already known. They are, however, to a large extent the subject of the Applicant's as yet unpublished patent applications and are obtainable by means of the processes described therein, for example by reacting

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4-alkylideneimino-lH-triazolinones of the formula (IX),

in which

R1, R14 and R15 have the meaning given above,

with halogenobenzene derivatives of the formula (III),

5 in which

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 R^3 , R^4 , R^5 , R^6 and R^7 have the meaning given above and Hal^1 represents halogen,

optionally in the presence of a diluent and optionally in the presence of a reaction auxiliary, in analogy to the implementation of process (a) according to the invention.

4-Alkylidenimino-1H-triazolinones of the formula (IX) are known or are obtainable by analogy with known processes

(cf. e.g. EP 294 666; EP 399 294).

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Suitable diluents for carrying out process (a) according to the invention are inert organic solvents. These include, in particular, aliphatic, alicyclic or aromatic, optionally halogenated hydrocarbons such as, for example, benzine. benzene. toluene, xylene, chlorobenzene. dichlorobenzene, petroleum ether, hexane, cyclohexane, dichloromethane, chloroform or carbon tetrachloride; ethers such as diethyl ether, disopropyl ether, dioxane, tetrahydrofuran or ethylene glycol dimethyl or diethyl ether; ketones such as acetone, butanone or methyl isobutyl ketone; nitriles such as acetonitrile, propionitrile or benzonitrile; amides such as N,N-dimethylformamide, N.N-dimethylacetamide, N-methylformanilide, Nmethylpyrrolidone or hexamethylphosphoric triamide or esters such as methyl acetate or ethyl acetate.

Process (a) according to the invention is preferably carried out in the presence of a suitable reaction auxiliary. Suitable such auxiliaries are all conventional inorganic or organic bases. These include, for example, alkaline earth metal or alkali metal hydroxides such as sodium hydroxide, calcium hydroxide, potassium hydroxide or else ammonium hydroxide, alkali metal carbonates such as sodium carbonate, potassium carbonate, potassium hydrogen carbonate, sodium hydrogen carbonate or ammonium carbonate, alkali metal or alkaline earth metal acetates such as sodium acetate, potassium acetate, calcium acetate or ammonium acetate, and tertiary amines such as trimethylamine, triethylamine, tributylamine,

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N.N-dimethylaniline, pyridine, piperidine, N-methylpiperidine, N, N-dimethylaminopyridine, diazabicyclooctane (DABCO), diazabicyclononene (DBN) or diazabicycloundecene (DBU).

- 5 When carrying out process (a) according to the invention, the reaction temperatures can be varied within a relatively wide range. It is in general carried out at temperatures of between 0°C and +180°C, preferably at temperatures of between +20°C and +120°C.
- 10 Process (a) according to the invention is usually carried out under atmospheric pressure. However, it is also possible to work under increased or reduced pressure.

To carry out process (a) according to the invention requires the use, per mole of 1H-triazolinone of the formula (II), of in general from 1.0 to 3.0 mol, prefer-15 ably from 1.0 to 1.5 mol, of halogenobenzene derivative of the formula (III) and optionally from 1.0 to 3.0 mol, preferably from 1.0 to 1.5 mol, of base as reaction auxiliary. The reaction procedure, work-up and isolation of the reaction products are carried out by known processes which are generally conventional (compare also the Preparation Examples).

Suitable diluents for carrying out process (b) according to the invention are inert organic solvents. It is preferred to use the solvents listed in the description of the implementation of process (a) according to the

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invention.

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Process (b) according to the invention is preferably carried out in the presence of a suitable reaction auxiliary. Suitable such auxiliaries are all conventional inorganic or organic bases. These include, for example, alkaline earth metal or alkali metal hydrides, hydroxides, amides, alcoholates, acetates, carbonates or hydrogen carbonates such as, for example, sodium hydride, sodium amide, sodium methylate, sodium ethylate, potassium tert-butylate, sodium hydroxide, potassium hydroxide, ammonium hydroxide, sodium acetate, potassium acetate, calcium acetate, ammonium acetate, sodium carbonate, potassium carbonate, potassium hydrogen carbonate, sodium hydrogen carbonate or ammonium carbonate and tertiary amines such as trimethylamine, triethylamine, tributylamine, N,N-dimethylaniline, pyridine, Nmethylpiperidine, N,N-dimethylaminopyridine, diazabicyclooctane (DABCO), diazabicyclononene (DBN) or diazabicycloundecene (DBU).

- When carrying out process (b) according to the invention, the reaction temperatures can be varied within a relatively wide range. It is generally carried out at temperatures of between -20°C and +150°C, preferably at temperatures of between 0°C and +120°C.
- 25 Process (b) according to the invention is usually carried out under atmospheric pressure. However, it is also possible to work under increased or reduced pressure.

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To carry out process (b) according to the invention requires the use, per mole of substituted triazolinone of the formula (Ia), of in general from 1.0 to 3.0 mol, preferably from 1.0 to 1.5 mol, of nucleophile of the formula (IV) and optionally from 0.1 to 3.0 mol, preferably from 1.0 to 1.5 mol, of base as reaction auxiliary. The reaction procedure, work-up and isolation of the reaction products are carried out by known processes which are generally conventional.

Suitable diluents for carrying out process (c) according 10 to the invention are inert organic solvents. These include, in particular, aliphatic, alicyclic or aromatic, optionally halogenated hydrocarbons such as, for example, benzine, benzene, toluene, xylene, chlorobenzene, dichlorobenzene, petroleum ether, hexane, cyclohexane, 15 dichloromethane, chloroform, carbon tetrachloride; ethers such as diethyl ether, disopropyl ether, dioxane, tetrahydrofuran or ethylene glycol dimethyl or diethyl ether; nitriles such as acetonitrile, propionitrile or 20 benzonitrile; amides, such as N, N-dimethylformamide, N, Ndimethylacetamide, N-methylformanilide, N-methylpyrrolidone or hexamethylphosphoric triamide; esters such as methyl acetate or ethyl acetate or sulphoxides such as dimethyl sulphoxide.

Process (c) according to the invention can optionally also be carried out in a two-phase system such as, for example, water/toluene or water/dichloromethane, optionally in the presence of a suitable phase-transfer

catalyst. Examples of such catalysts which may be mentioned are: tetrabutylammonium iodide, tetrabutylammonium bromide, tetrabutylammonium chloride, tributyl-methyl-phosphonium bromide, trimethyl-C₁₃/C₁₅-alkylammonium chloride, trimethyl-C₁₃/C₁₅-alkylammonium bromide, dibenzyl-dimethyl-ammonium methyl sulphate, dimethyl-C₁₂/C₁₄-alkyl-benzylammonium chloride, dimethyl-C₁₂/C₁₄-alkyl-benzylammonium bromide, tetrabutylammonium hydroxide, triethylbenzylammonium chloride, methyltrioctylammonium chloride, trimethylbenzylammonium chloride, 15-crown-5, 18-crown-6 or tris-[2-(2-methoxyethoxy)-ethyl]-amine.

Process (c) according to the invention is preferably carried out in the presence of a suitable reaction auxiliary. Suitable such auxiliaries are all conventional inorganic or organic bases. These include, for example, alkaline earth metal or alkali metal hydrides, hydroxides, amides, alcoholates, acetates, carbonates or hydrogen carbonates such as, for example, sodium hydride, sodium amide, sodium methylate, sodium ethylate, potassium tert-butylate, sodium hydroxide, potassium hydroxide, ammonium hydroxide, sodium acetate, potassium acetate, calcium acetate, ammonium acetate, carbonate, potassium carbonate, potassium hydrogen carbonate, sodium hydrogen carbonate or ammonium carbonate and tertiary amines such as trimethylamine, triethylamine, tributylamine, N,N-dimethylaniline, pyridine, Nmethylpiperidine, N,N-dimethylaminopyridine, diazabicyclooctane (DABCO), diazabicyclononene (DBN) or diazabicycloundecene (DBU).

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When carrying out process (c) according to the invention, the reaction temperatures can be varied within a relatively wide range. It is in general carried out at temperatures of between -20°C and +150°C, preferably at temperatures of between 0°C and +120°C.

Process (c) according to the invention is usually carried out under atmospheric pressure. However, it is also possible to work under increased or reduced pressure.

To carry out process (c) according to the invention requires the use, per mole of substituted triazolinone of the formula (V), of in general from 1.0 to 3.0 mol, preferably from 1.0 to 2.0 mol, of alkylating, acylating or sulphonylating agent of the formula (VI) and optionally from 1.0 to 3.0 mol, preferably from 1.0 to 2.0 mol, of base as reaction auxiliary.

The reaction procedure, work-up and isolation of the reaction products are carried out in both cases by known processes which are generally conventional.

Suitable reducing agents for carrying out process (d)

according to the invention are conventional reducing
agents. It is particularly preferred to use complex
hydrides such as, for example, lithium aluminium hydride
or sodium borohydride.

Suitable diluents for carrying out process (d) according to the invention are, depending on the reducing agent used, conventional organic or inorganic solvents. The

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preferred diluents used are alcohols such as methanol, ethanol, propanol or butanol, ether alcohols such as methoxyethanol or ethoxyethanol, ethers such as diethyl ether, diisopropyl ether, dioxane or tetrahydrofuran, and their mixtures with water, or water alone.

When carrying out process (d) according to the invention, the reaction temperatures can be varied within a relatively wide range. It is generally carried out at temperatures of between -20°C and +100°C, preferably at temperatures of between 0°C and +80°C.

Process (d) according to the invention is usually carried out under atmospheric pressure. However, it is also possible to work under increased or reduced pressure.

To carry out process (d) according to the invention requires the use, per mole of 4-alkylidenimino-triazolin-one of the formula (VII), of in general from 0.5 to 5.0 mol, preferably from 1.0 to 3.0 mol, of reducing agent. The reaction procedure, work-up and isolation of the reaction products are carried out by known processes which are generally conventional.

The purification of the end products of the formula (I) is carried out by means of known methods, for example by column chromatography or by recrystallization.

Characterization is made via the melting point or, in the case of non-crystallizing compounds, by means of proton nuclear magnetic resonance spectroscopy (1H-NMR).

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The active compounds according to the invention can be used as defoliants, desiccants, agents for destroying broad-leaved plants and, especially, as weed-killers. By weeds, in the broadest sense, there are to be understood all plants which grow in locations where they are undesired. Whether the substances according to the invention act as total or selective herbicides depends essentially on the amount used.

The active compounds according to the invention can be used, for example, in connection with the following plants:

Dicotyledon weeds of the genera: Sinapis, Lepidium, Galium, Stellaria, Matricaria, Anthemis, Galinsoga, Chenopodium, Urtica, Senecio, Amaranthus, Portulaca, Kanthium, Convolvulus, Ipomosa, Polygonum, Sesbania, Ambrosia, Cirsium, Carduus, Sonchus, Solanum, Rorippa, Rotala, Lindernia, Lamium, Veronica, Abutilon, Emex, Datura, Viola, Galeopsis, Papaver Centaurea, Trifolium, Ranunculus and Taraxacum.

20 <u>Dicotyledon cultures of the genera:</u> Gossypium, Glycine, Beta, Daucus, Phaseolus, Pisum, Solanum, Linum, Ipomoea, Vicia, Nicotiana, Lycopersicon, Arachis, Brassica, Lactuca, Cucumis and Cucurbita.

Monocotyledon weeds of the genera: Echinochloa, Setaria,
25 Panicum, Digitaria, Phlaum, Poa, Festuca, Eleusine,
Brachiaria, Lolium, Bromus, Avena, Cyperus, Sorghum,
Agropyron, Cynodon, Monochoria, Fimbristylis, Sagittaria,
Eleocharis, Scirpus, Paspalum, Ischaemum, Sphenoclea,
Dactyloctenium, Agrostis, Alopecurus and Apera.

30 Monocotyledon cultures of the genera: Oryza, Zea,

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Triticum, Hordeum, Avena, Secale, Sorghum, Panicum, Saccharum, Ananas, Asparagus and Allium.

However, the use of the active compounds according to the invention is in no way restricted to these genera, but also extends in the same manner to other plants.

The compounds are suitable, depending on the concentration, for the total combating of weeds, for example on industrial terrain and rail tracks and on paths and squares with or without trees planted. Equally, the compounds can be employed for combating weeds in perennial cultures, for example afforestations, decorative tree plantings, orchards, vineyards, citrus groves, nut orchards, banana plantations, coffee plantations, tea plantations, rubber plantations, oil palm plantations, cocoa plantations, soft fruit plantings and hopfields, in lawns, turf and pasture-land, and for the selective combating of weeds in annual cultures.

In this context, the active compounds according to the invention can be employed with particularly good success for combating dicotyledon weeds in mono- and dicotyledon cultures such as, for example, soya, sunflower or barley. In addition, the active compounds according to the invention also possess, at corresponding application rates, fungicidal activity and can be employed for combating diseases in rice, such as, for example, against the causative organism of rice blast disease (Pyricularia oryzae).

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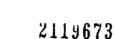
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Depending on their particular physical and/or chemical properties, the active compounds can be converted to the customary formulations, such as solutions, emulsions, suspensions, powders, foams, pastes, granules, aerosols, natural and synthetic materials impregnated with active compound, very fine capsules in polymeric substances and in coating compositions for seed, and furthermore in formulations used with burning equipment, such as fumigating cartridges, fumigating cans, fumigating coils and the like, as well as ULV cold mist and warm mist formulations.

These formulations are produced in a known manner, for example by mixing the active compounds with extenders, that is, liquid solvents, liquefied gases under pressure, and/or solid carriers, optionally with the use of surface-active agents, that is, emulsifying agents and/or dispersing agents, and/or foam-forming agents. In the case of the use of water as an extender, organic solvents can, for example, also be used as auxiliary solvents. As liquid solvents, there are suitable in the main: aromatics, such as xylene, toluene or alkylnaphthalenes. chlorinated aromatics or chlorinated aliphatic hydrocarbons, such as chlorobenzenes, chloroethylenes or methylene chloride, aliphatic hydrocarbons, such as cyclohexane or paraffins, for example mineral oil fractions, alcohols, such as butanol or glycol as well as their ethers and esters, ketones, such as acetone, methyl ethyl ketone, methyl isobutyl ketone or cyclohexanone, strongly polar solvents, such as dimethylformamide and





dimethyl sulphoxide, as well as water; by liquefied gaseous extenders or carriers are meant those liquids which are gaseous at ambient temperature and under atmospheric pressure, for example aerosol propellants, such as halogenated hydrocarbons as well as butane, propane, nitrogen and carbon dioxide; as solid carriers there are suitable: for example ground natural minerals, such as kaolins, clays, talc, chalk, quartz, attapulgite, montmorillonite or diatomaceous earth, and ground synthetic minerals, such as highly disperse silica, alumina and silicates; as solid carriers for granules there are suitable: for example crushed and fractionated natural rocks such as calcite, marble, pumice, sepiolite and dolomite, as well as synthetic granules of inorganic and organic meals, and granules of organic material such as sawdust, coconut shells, maize cobs and tobacco stalks; as emulsifying and/or foam-forming agents there are suitable: for example non-ionic and anionic emulsifiers, such as polyoxyethylene fatty acid esters, polyoxyethylene fatty alcohol ethers, for example alkylaryl polyglycol ethers, alkylsulphonates, alkyl sulphates, arylsulphonates as well as albumen hydrolysis products; as dispersing agents there are suitable: for example lignin-sulphite waste liquors and methylcellulose.

Adhesives such as carboxymethylcellulose and natural and synthetic polymers in the form of powders, granules or latices, such as gum arabic, polyvinyl alcohol and polyvinyl acetate, as well as natural phospholipids, such as cephalins and lecithins, and synthetic phospholipids,

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can be used in the formulations. Other additives can be mineral and vegetable oils.

It is possible to use colorants such as inorganic pigments, for example iron oxide, titanium oxide and Prussian Blue, and organic dyestuffs, such as alizarin dyestuffs, azo dyestuffs and metal phthalocyanine dyestuffs, and trace nutrients such as salts of iron, manganese, boron, copper, cobalt, molybdenum and tin.

The formulations in general contain between 0.1 and 95
10 per cent by weight of active compound, preferably between
0.5 and 90%.

For controlling weeds, the active compounds according to the invention, as such or in the form of their formulations, can also be used as mixtures with known herbicides, finished formulations or tank mixes being possible.

Known herbicides are suitable for the mixtures, for example anilides, such as for example diflufenican and propanil; arylcarboxylic acids, for dichloropicolinic acid, dicamba or picloram; aryloxyalkanoic acids, for example 2,4-D, 2,4-DB, 2,4-DP, fluroxypyr, MCPA, MCPP and triclopyr; aryloxy-phenoxyalkanoic acid esters, for example diclofop-methyl, fenoxaprop-ethyl, fluazifop-butyl, haloxyfop-methyl and quizalofop-ethyl; azinones, for example chloridazon and norflurazon; carbamates, for example chlorpropham, desmedipham, phenmedipham and propham;





chloroacetanilides, for example alachlor, acetochlor, butachlor, metazachlor, metolachlor, pretilachlor and propachlor; dinitroanilines, for example oryzalin, pendimethalin and trifluralin; diphenyl ethers, for example acifluorfen, bifenox, fluoroglycofen, fomesafen, halosafen, lactofen and oxyfluorfen; ureas, for example chlortoluron, diuron, fluometuron, isoproturon, linuron and methabenzthiazuron; hydroxylamines, for example alloxydim. clethodim, cycloxydim, sethoxydim tralkoxydim; imidazolinones, for example imazethapyr, imazamethabenz, imazapyr and imazaquin; nitriles, for example bromoxynil, dichlobenil and ioxynil; oxyacetamides, for example mefenacet; sulphonylureas, for example amidosulfuron, bensulfuron-methyl, chlorimuronethyl, chlorsulfuron, cinosulfuron, metsulfuron-methyl, nicosulfuron, primisulfuron, pyrazosulfuron-ethyl, thifensulfuron-methyl, triasulfuron and tribenuronmethyl; thiocarbamates, for example butylate, cycloate, diallate, EPTC, esprocarb, molinate, prosulfocarb, thiobencarb and triallate; triazines, for example atrazine, cyanazine, simazine, simetryne, terbutryne and terbutylazine; triazinones, for example hexazinone, metamitron and metribuzin; and others, for example aminotriazole, benfuresate, bentazone, cinmethylin. clomazone, clopyralid, difenzoquat, dithiopyr, ethofumesate, fluorochloridone, glufosinate, glyphosate, isoxaben, pyridate, quinchlorac, quinmerac, sulphosate and tridiphane.

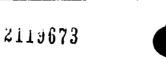
Mixtures with other known active compounds, such as

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fungicides, insecticides, acaricides, nematicides, bird repellants, plant nutrients and agents which improve soil structure, are also possible.

- The active compounds can be used as such, in the form of their formulations or in the use forms prepared therefrom by further dilution, such as ready-to-use solutions, suspensions, emulsions, powders, pastes and granules. They are used in the customary manner, for example by watering, spraying, atomizing or scattering.
- 10 The active compounds according to the invention can be applied either before or after emergence of the plants. They can also be incorporated into the soil before sowing.
- The amount of active compound used can vary within a 15 relatively wide range. It depends essentially on the nature of the desired effect. In general, the application rates are between 0.01 and 10 kg of active compound per hectare of soil surface, preferably between 0.05 and 5 kg per hectare.
- 20 The preparation and use of the active compounds according to the invention can be seen from the following examples.





Preparation Examples:

Example 1:

(Process a)

83 g (0.06 mol) of potassium carbonate are added to 71 g (0.5 mol) of 4-dimethylamino-3-methyl-1H-1,2,4-triazolin-5-one (cf. e.g. EP 422 469) and 78.5 g (0.5 mol) of 2,4,5-trifluorobenzonitrile (cf. e.g. EP 191 181) in 400 ml of dimethyl sulphoxide at room temperature and the mixture is then stirred at 40°C to 50°C for two hours.

10 For working up, the cooled reaction mixture is filtered, the filtrate is concentrated in vacuo, the residue is stirred together with water, the precipitated solid is filtered off with suction, washed with water and dried.

111 g (80% of theory) of 1-(4-cyano-2,5-difluorophenyl)3-methyl-4-dimethylamino-1,2,4-triazolin-5-one are
obtained with a melting point of 116°C.

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Example 2:

(Process b)

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0.6 g (0.015 mol) of sodium hydride (60% in paraffin oil) is added at room temperature to 1.05 g (0.015 mol) of 3-butin-1-ol in 100 ml of acetonitrile, the mixture is stirred for 10 minutes at room temperature, then 2.12 g (0.008 mol) of 1-(4-cyano-2,5-difluorophenyl)-3-methyl-4-(N-methylamino)-1,2,4-triazolin-5-one are added and the mixture is stirred for a further 16 hours at room temperature. For working up, the reaction mixture is filtered, the filtrate is concentrated in vacuo, the residue is stirred together with water, and the precipitated solid is filtered off with suction, washed with water and dried.

1.96 g (78% of theory) of 1-(4-cyano-2-fluoro-5-but-1-in-3-yl-oxyphenyl)-3-methyl-4-(N-methylamino)-1,2,4-triazo-lin-5-one is obtained with a melting point of 184-185°C.

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Example 3:

(Process a)

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16.5 g (0.12 mol) of potassium carbonate are added to 12.8 g (0.1 mol) of 4-(N-methylamino)-3-methyl-1H-1,2,4-triazolin-5-one (cf. e.g. EP 399 294) and 15.7 g (0.1 mol) of 2,4,5-trifluorobenzonitrile (cf. e.g. EP 191 181) in 200 ml of dimethyl sulphoxide at room temperature, and the mixture is then stirred at 40°C to 50°C for three hours. For working up, the cooled reaction mixture is filtered, the filtrate is concentrated in vacuo, the residue is stirred together with water, and the precipitated solid is filtered off with suction, washed with water and dried.

12.8 g (48% of theory) of 1-(4-cyano-2,5-difluoro-15 phenyl)-3-methyl-4-(N-methylamino)-1,2,4-triazolin-5-one are obtained with a melting point of 128-131°C.

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Example 4:

(Process a)

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1.9 g (0.014 mol) of potassium carbonate is added to 1.9 g (0.012 mol) of 3-methyl-4-(N-isopropylamino)-1H-1,2,4-triazolin-5-one (preparation analogous to EP 399 294) and 1.9 g (0.012 mol) of 2,4,5-trifluorobenzonitrile (cf. e.g. EP 191 181) in 100 ml of dimethyl sulphoxide at room temperature, and the mixture is then stirred at room temperature for two hours and at 40-50°C for 1.5 hours. For working up, the cooled reaction mixture is placed in water, and the precipitated solid is filtered off with suction, washed with water and dried.

1.3 g (54.3% of theory) of 1-(4-cyano-2,5-difluoro-phenyl)-3-methyl-4-(N-isopropylamino)-1,2,4-triazolin-5-one is obtained with a melting point of 35-36°C.

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Preparation of the starting compounds:

Example III-1:

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220 g (1.06 mol) of 2,4,5-trichlorobenzonitrile are added with stirring at room temperature to 250 g (4.31 mol) of potassium fluoride in 400 ml of distilled tetramethylene sulphone, and the mixture is then stirred at 195°C to 200°C for 10 hours. For working up, the mixture is cooled, 500 ml of water are added, and the mixture is subjected to steam distillation. The organic fraction is taken up in dichloromethane, dried over sodium sulphate, concentrated in vacuo and distilled.

108 g (58% of theory) of 2,4-difluoro-5-chlorobenzonitrile are obtained with a boiling point of 105-107°C at 30 mbar and with a melting point of 48-50°C.

In a corresponding manner, and in accordance with the general instructions for the preparation, the following substituted triazolinones of the general formula (I) are obtained:

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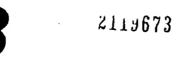
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Ex.No.	R	R^3 R^4 R^5	physical properties
7	H,C NH-CH,	NH-SO ₂ -CH ₃	m.p. >250°C
8	H ₃ C ₂ N(CH ₂) ₂	F CN	m.p. 81°C
9	H ₂ C ₂ N(CH ₂) ₂	CH, O-CH-CECH	¹ H-NMR ^{*)} : 1.73-1.75; 3.0; 4.92-5.0
10	H ₂ C ₂ N(CH ₂) ₂	O-CH ₂ -C=CH	¹ H-NMR ^{*)} : 2.6-2.7; 3.02; 4.85





Ex.No.



R³ R⁴

physical properties

11	H ₂ C ₂ N(CH ₃) ₂	(O-CH ₂ -CH ₂) ₂ -N(CH ₂) ₂	¹ H-NMR*):
	N /		2.30; 3.0;
	, MO	\	4.25-4.30
		F	





Ex.No.

$$R^{3}$$
 R^{4}
 R^{5}
 R^{6}

physical properties

14	H,C N(CH,)	NH-SO ₂ -C ₂ H ₃ m.p. 192-193°C
	N N	
	, A	<u></u>
15	ңс <u>мнс</u> ң	F ,0—CH₂-C≡CH m.p. 160-161°C
	N N	

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Ex.No.

R R R

R⁷ R⁶

physical properties

17 H,C NH-CH, (O-CH₂-CH₂)₂-OCH, 1_{H-NMR}*); 2.35; 2.76-2.79; 3.4; 4.25-4.3

18 H,C NH-CH, O-CH,-CH=CH, m.p. 137-138°C

19 H₂C NH-CH₃ m.p. 168-169°C





Ex.No.	R ¹ R ² X	R^3 R^4 R^6	, physical R properties
20 H ₂ C	NHCH,	O-CH ₂	m.p. 158-160°C



Ex.No.	R ¹ R ² N X	$ \begin{array}{c} R^3 \\ R^4 \\ R^5 \end{array} $	physical properties
	_	NTV 00 011	

23	N(CH ₂) ₂	NH-SO ₂ -C ₂ H ₃	m.p. 185-187°C	
24	1	CH,	m.p. 106°C	

Ex.No		R ³ R ⁴ physical properties
26	H ₂ C ₂ NH-CH ₃	NH-SO ₂ -C ₂ H ₃ m.p. 170°C
27	H ₂ C ₂ NH-CH ₃	NH-SO ₂ -n-C ₄ H ₉ m.p. 183-185°C
28	H ₂ C ₂ NH-CH ₃	NH-SO ₂ -CH ₃ m.p. 176°C
29	H ₂ C ₂ NH-CH ₃	O—CH ₂ -CH ₂ -CH ₂ m.p. 100-102°C



Ex.No.	R ¹ R ² X	$ \begin{array}{c} R^{3} \\ R^{5} \end{array} $ $ \begin{array}{c} R^{6} \end{array} $	physical properties
30	H,C,-i NH-CH,	F cn	.p. 112-113°C
31	NH-CH,	F m.	р. 136-138°С
32	H,C ₃ -i NH-CH,	O-CH-CECH m.	p. 121-123℃



Ex.No.	R R X	R ³ R ⁴ physical properties
33	H ₂ C ₂ -i NH-CH ₃	NH-SO ₂ -CH ₃ m.p. 168-170°C
34	NH-CH ₃	NH-SO ₂ -CH ₄ m.p. 155-157°C
35	H,C NH-CH,	NH-SO ₂ -C ₂ H ₃ m.p. 202-204°C



Ex.No.	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	\mathbb{R}^{3} \mathbb{R}^{4} \mathbb{R}^{6}	of physical properties
36	H,C NH-CH,	NH-i-C,H,	m.p. 188-190°C
37	H,C NH-CH,	NH-CH ₂ -CH ₂ -CH ₂	m.p. 158-160°С
38	H,C NH-CH,	-CN	m.p. 117-119°C
39	H,C NHCH,	NH-SO ₂ -n-C ₄ H ₅ —CN	m.p. 128-130°C



Ex.No	R R X	R ¹ R ⁴ physical properties
40	H ₃ C N(CH ₃) ₂	NH-SO ₂ -a-C ₄ H ₉ m.p. 146-148°C ———————————————————————————————————
41	H,C NH-CH,	NH-SO ₂ -n-C ₄ H _g m.p. 106-108°C
42	H ₂ C NH-CH ₃	m.p. 146-148°C



Ex.No.	R R X	R ³ R ⁴ physical properties
43	H,C NH-CH,	F m.p. 125-126°C CF ₃
44	H,C NH-CH,	CI m.p. 107-108°C
45	H,C NH-CH	CI F m.p. 115-118°C

Ex.No.	R R	ex 🔷	P physical properties
46	H,C N(CH ₂) ₂	`ocH-cH*o- cH	m.p. 85-87°С -СЦ
	N N O	-CN	





Ex.No.	R R R X	R ³ R ⁴	ophysical properties
50	H ₂ C N(CH ₂) ₂	NH-SO ₂ -n-C _e H _g —CN	m.p. 151-152°C
51	H,C NH-CH,	CH ₃	m.p. 178-179°C
52	H,C NHCH,		m.p. 227-228°C
53	H,C N(CH ₂) ₂	O—CH ₂ -C≡CH	m.p. 87-89°C



Ex.No.	R ¹ R ² N X	R ³ R ⁴ physical properties
54	H,C NH-CH,	CI NH-SO ₂ -CH ₃ m.p. 187-188°C ———————————————————————————————————
55	HCS NH-CH	NH-SO ₂ -C ₂ H ₃ m.p. 160-161°C ———————————————————————————————————
56	H ₂ C N(CH ₂) ₂	O-CH ₂ -CH ₂ -SC ₂ H ₃ m.p. 95-97°C



Ex.No	R ¹ R ²	R ³ R ⁴ physical properties
57	H,C N(CH,)2	O-CH ₂ -CF ₂ -CF ₃ m.p. 113°C
	N NO	F
58	H,C N(CH,)2	CH, m.p. 212-214°C
	N N O	H ₂ C————————————————————————————————————



Ex.No	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	R ³ R ⁴	physical properties
59	F ₃ C N(CO-CH ₃) ₂ N N O	F CN	¹ H-NMR ^{*)} : 2.52; 7.55-7.65
60	F,C NH-CO-CF,	F CN	¹ H-NMR ^{*)} : 7.55-7.60
61	H ₂ C N(CH ₂) ₂	N(CH ₃)-SO ₂ -n-C ₄ H ₃ m.	.p. 121-123°C
62	H,CS NH-CH,	F m.	p. 150-151°C





Ex.No.	R ¹ R ² N X	R ³ R ⁴ physical properties
63	F,C NH-CO-CF,	NH-SO ₂ -C ₂ H, 1 _{H-NMR} *): 1.45; 3.2-3.25; 7.7; 7.95-7.98
64	H ₂ C NH-i-C ₃ H ₂	NH-SO ₂ -C ₂ H ₅ 1 _{H-NMR} *): 1.1-1.12; 2.3; 3.65-3.75; 4.58
65	H,C N(CH,)2	m.p. 130°C





Ex. No.	R ¹ R ²	R^3 R^4 R^5 R^7 R^6	Physical properties
66	H ₃ C NHCH ₃	CI F	m.p. 101°C
67	H ₃ C N(CH ₃) ₂	O-I-C ₃ H ₇ ———————————————————————————————————	¹ H-NMR*): 1.40-1.42; 2.3; 3.0; 4.6-4.7
68	CH ₃ NHCH ₃	SC ₂ H ₅ CN	m.p. 117-119°C
69	CH ₃ NHCH ₃	NH ₂ NO ₂	m.p. 151-152°C
70	CH ₃ S NH-CO-CF ₃	F_CN	m.p. 84-86°C
71	CH ₃ S NH-CO-CF ₃	NH-SO ₂ -C ₂ H ₅ —CN	m.p. 137-138°C



Ex. No.	R ¹ R ² N X	R^3 R^4 R^5 R^7 R^6	Physical properties
72	N NHCH ₃	CI CF ₃	m.p. 117-119°C
73	NHCH ³	F CN F	m.p. 120-122°C
74	CH ₃ NHCH ₃	OC₂H₅ CN	m.p. 161°C
75	CH ₃ N(CH ₃)₂ N(CH ₃)₂	NH-SO ₂ -CH ₃	m.p. 149°C
76	CH ₃ NHCH ₃	NH-SO ₂ -CH ₃ —CN	m.p. 143°C
77	CH ₃ N(CH ₃) ₂	OC ₂ H ₅ CN	m.p. 89°C

Ex. No.	R ¹ R ² N X	R^3 R^4 R^5 R^7 R^6	Physical properties
78	N(CH ₃) ₂	F———CN	m.p. 103°C
79	CF ₃ N HCH ₃	F_CN	m.p. 145°C

*) The ¹H-NMR spectra were recorded in deuterochloroform (CDCl₃) using tetramethylsilane (TMS) as the internal standard. The value given is the chemical shift δ in ppm.

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Application Examples:

In the following Application Example, the compound listed below was employed as comparison substance:

3-Methyl-4-propargyl-1-(2,5-difluoro-4-cyano-phenyl)-1,2,4-triazolin-5-one

(known from DE 38 39 480)





Example A:

Pre-emergence test

Solvent:

5

10

15

20

25

5 parts by weight of acetone

Emulsifier:

1 part by weight of alkylaryl polyglycol

ether

To produce a suitable preparation of active compound, one part by weight of active compound is mixed with the stated amount of solvent, the stated amount of emulsifier is added and the concentrate is diluted with water to the desired concentration.

Seeds of the test plants are sown in normal soil and, after 24 hours, watered with the preparation of the active compound. It is expedient to keep constant the amount of water per unit area. The concentration of an active compound in the preparation is of no importance, only the amount of active compound applied per unit area being decisive. After three weeks, the degree of damage to the plants is rated in % damage in comparison to the development of the untreated control.

The figures denote:

0% = no action (like untreated control)
100% = total destruction

In this test, for example the compounds according to Preparation Examples 5 and 6 exhibit a distinctly superior activity at a rate of 250 g/ha compared to the prior art, in cultures like soy-bean (0-30%), sunflowers (0%), barley (0-100%) against weeds like abuthilon (95-100%), chenopodium (100%), galium (80-95%), matricaria (95-100%) and solanum (95-100%) although the prior art has been applicated at a rate of 500 g/ha.



THE EMBODIMENTS OF THE INVENTION IN WHICH AN EXCLUSIVE PROPERTY OR PRIVILEGE IS CLAIMED ARE DEFINED AS FOLLOWS:

1. A substituted 1-aryltriazolinone of the general
formula (I):

in which

R¹ represents hydrogen, alkyl, halogenalkyl, alkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl or cycloalkyl,

 R^2 represents a radical of the formula $-NR^8R^9$,

 ${\mbox{R}}^3$, ${\mbox{R}}^6$ and ${\mbox{R}}^7$ independently of one another in each case represents hydrogen, halogen, amino or nitro,

 $\begin{array}{c} {\rm R}^4 \ \ {\rm represents} \ \ {\rm hydrogen}, \ \ {\rm halogen}, \ \ {\rm cyano} \ \ {\rm or} \ \ {\rm nitro}, \ \ {\rm or} \\ {\rm one} \ \ {\rm of} \ \ {\rm the} \ \ {\rm radicals} \ \ -{\rm R}^{10}, \ \ -{\rm O-R}^{10}, \ \ -{\rm S-R}^{10}, \ \ -{\rm S(0)-R}^{10}, \ \ -{\rm So}_2{\rm -R}^{10}, \\ {\rm -SO}_2{\rm -OR}^{10}, \ \ -{\rm SO}_2{\rm -NR}^{11}{\rm R}^{10}, \ \ -{\rm CO-OR}^{10}, \ \ -{\rm CO-NR}^{11}{\rm R}^{10}, \ \ -{\rm O-SO}_2{\rm -R}^{10}, \\ {\rm -N(R}^{11}){\rm -SO}_2{\rm -R}^{10}, \ \ -{\rm NR}^{11}{\rm R}^{10}, \ \ -{\rm NH-P(O)} \ ({\rm R}^{11}) \ ({\rm OR}^{10}) \ \ {\rm or} \\ {\rm -NH-P(O)} \ \ ({\rm OR}^{11}) \ \ ({\rm OR}^{10}), \end{array}$

 ${ t R}^{ extsf{5}}$ represents nitro, cyano, halogen or halogenoalkyl, and

X represents oxygen or sulphur, where

 $\rm R^8$ represents hydrogen, alkyl, halogenoalkyl, a radical of the formula -CO-R¹² or a radical of the formula -S(O) $_{\rm n}$ -R¹², $\rm R^9$ represents alkyl, halogenoalkyl, a radical of the



formula $-CO-R^{12}$ or a radical of the formula $-S(0)_n-R^{12}$, R^{10} represents hydrogen or represents in each case optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl, aryl,

optionally substituted alkyl, alkenyl, alkinyl, cycarylalkyl or heterocyclyl,

R¹¹ represents hydrogen or represents in each case optionally substituted alkyl, alkenyl, alkinyl, cycloalkyl, arylalkyl or aryl,

R¹² represents in each case optionally substituted alkyl, cycloalkyl, arylalkyl, aryl or heterocyclyl, and n represents a number 0, 1 or 2.

A substituted 1-aryltriazolinone of the general formula
 according to claim 1, characterized in that

R¹ represents hydrogen or represents in each case straight-chain or branched alkyl, alkoxy, alkylthic or alkyl-sulphonyl having in each case from 1 to 8 carbon atoms, furthermore represents straight-chain or branched halogenoalkyl having from 1 to 8 carbon atoms and from 1 to 17 identical or different halogen atoms, or represents cycloalkyl having from 3 to 8 carbon atoms,

 ${
m R}^2$ represents a radical of the formula $-{
m NR}^8{
m R}^9$, ${
m R}^3$, ${
m R}^6$ and ${
m R}^7$ independently of one another in each case represent hydrogen, fluorine, chlorine, bromine, iodine, amino or nitro,

 ${\rm R}^4$ represents hydrogen, fluorine, chlorine, bromine, iodine, cyano or nitro, or represents one of the radicals $-{\rm R}^{10}$, $-{\rm O-R}^{10}$, $-{\rm S-R}^{10}$, $-{\rm S}({\rm O})-{\rm R}^{10}$, $-{\rm SO}_2-{\rm R}^{10}$, $-{\rm SO}_2-{\rm OR}^{10}$, $-{\rm SO}_2-{\rm NR}^{11}{\rm R}^{10}$,



 $-\text{CO-OR}^{10}$, $-\text{CO-NR}^{11}\text{R}^{10}$, $-\text{O-SO}_2-\text{R}^{10}$, $-\text{N}(\text{R}^{11})-\text{SO}_2-\text{R}^{10}$, $-\text{NR}^{11}\text{R}^{10}$, $-\text{NH-P}(O)(\text{R}^{11})(\text{OR}^{10})$ or $-\text{NH-P}(O)(\text{OR}^{11})(\text{OR}^{10})$,

R⁵ represents nitro, cyano, fluorine, chlorine, bromine, iodine or represents straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms and

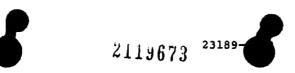
X represents oxygen or sulphur, where

 R^8 represents hydrogen, straight-chain or branched alkyl having from 1 to 8 carbon atoms or straight-chain or branched halogenoalkyl having from 1 to 8 carbon atoms and from 1 to 17 identical or different halogen atoms, and furthermore represents a radical of the formula $-CO-R^{12}$ or a radical of the formula $-S(O)_n-R^{12}$,

 ${
m R}^9$ represents straight-chain or branched alkyl having from 1 to 8 carbon atoms or straight-chain or branched halogenoalkyl having from 1 to 8 carbon atoms and from 1 to 17 identical or different halogen atoms, and furthermore represents a radical of the formula $-{
m CO-R}^{12}$ or a radical of the formula $-{
m S(O)}_{
m n}-{
m R}^{12}$,

R¹⁰ represents hydrogen,

R¹⁰ furthermore represents straight-chain or branched alkyl having from 1 to 14 carbon atoms which is optionally substituted once or more than once by identical or different substituents of halogen, cyano, carboxyl, carbamoyl, in each case straight-chain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, N-alkylamino-carbonyl, N,N-dialkylaminocarbonyl or alkylsulphonylaminocarbonyl having in each case from 1 to 8 carbon atoms in the individual



alkyl moieties, or heterocyclyl, the heterocyclyl radical being a five- to seven-membered optionally benzo-fused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur,

R¹⁰ furthermore represents alkenyl or alkinyl having in each case from 2 to 8 carbon atoms, which are optionally substituted once or more than once by identical or different halogens,

R¹⁰ furthermore represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once or more than once by identical or different substituents of halogen and/or straight-chain or branched alkyl having from 1 to 4 carbon atoms,

R¹⁰ furthermore represents arylalkyl or aryl having in each case from 6 to 10 carbon atoms in the aryl moiety and optionally from 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the aryl moiety once or more than once by identical or different substituents, or represents a saturated or unsaturated, five- to seven-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur, which is optionally substituted once or more than once by identical or different substituents and/or is benzo-fused, wherein substituents of the aryl and/or heterocyclyl are halogen, cyano, nitro, amino, N-acetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 6 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl



or halogenoalkylsulphonyl having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoxy-iminoalkyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once or more than once by identical or different substituents of halogen and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 6 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms,

R¹¹ represents hydrogen,

R¹¹ furthermore represents straight-chain or branched alkyl having from 1 to 14 carbon atoms which is optionally substituted once or more than once by identical or different substituents of fluorine, chlorine, bromine, iodine, cyano, carboxyl, carbamoyl, in each case straight-chain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, N-alkylaminocarbonyl, N,N-dialkylaminocarbonyl or alkylsulphonylaminocarbonyl having in each case from 1 to 8 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl radical being a five- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur,

R¹¹ furthermore represents alkenyl or alkinyl having in each case from 2 to 8 carbon atoms, which are optionally



substituted once or more than once by identical or different halogen atoms of fluorine, chlorine, bromine and/or iodine,

R¹¹ furthermore represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once or more than once by identical or different substituents of fluorine, chlorine, bromine, iodine and/or straight-chain or branched alkyl having from 1 to 4 carbon atoms,

R¹¹ furthermore represents arylalkyl or aryl having in each case from 6 to 10 carbon atoms in the aryl moiety and optionally from 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the aryl moiety once or more than once by identical or different substituents, wherein substituents of the aryl are in each case halogen, cyano, nitro, amino, N-acetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 6 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoxyiminoalkyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once or more than once by identical or different substituents of halogen and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 6 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 6 carbon atoms and from 1 to 13 identical

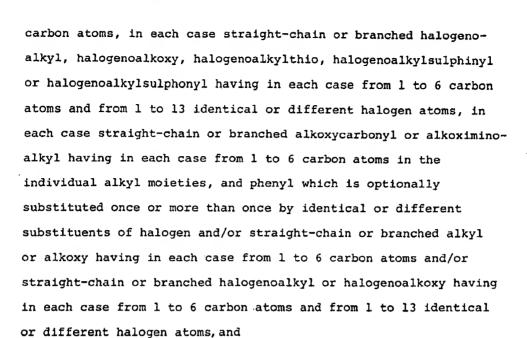


or different halogen atoms,

R¹² represents straight-chain or branched alkyl having from 1 to 8 carbon atoms which is optionally substituted once or more than once by identical or different substituents of fluorine, chlorine, bromine, iodine, cycloalkyl having from 3 to 8 carbon atoms or heterocyclyl, the heterocyclyl radical being a five- to seven-membered optionally benzo-fused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur,

R¹² furthermore represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once or more than once by identical or different substituents of fluorine. chlorine, bromine, iodine and/or straight-chain or branched alkyl having from 1 to 4 carbon atoms,

R¹² furthermore represents arylalkyl or aryl having in each case from 6 to 10 carbon atoms in the aryl moiety and optionally from 1 to 4 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the aryl moiety once or more than once by identical or different substituents, or represents a saturated or unsaturated, five- to seven-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur, which is optionally substituted once or more than once by identical or different substituents, wherein the substituents of aryl or heterocyclyl are in each case halogen, cyano, nitro, amino, N-acetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 6



n represents a number 0, 1 or 2.

A substituted 1-aryltriazolinone of the general formula
 according to claim 1, characterized in that

R¹ represents hydrogen or in each case straight-chain or branched alkyl, alkoxy, alkylthio or alkylsulphonyl having in each case from 1 to 6 carbon atoms, or furthermore represents straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms of fluorine, chlorine or bromine, or represents cycloalkyl having from 3 to 7 carbon atoms,

 R^2 represents a radical of the formula $-NR^8R^9$,

 ${\rm R}^3$, ${\rm R}^6$ and ${\rm R}^7$ independently of one another in each case represent hydrogen, fluorine, chlorine, bromine, amino or nitro,

R4 represents hydrogen, fluorine, chlorine, bromine,

cyano or nitro, or represents one of the radicals $-R^{10}$, $-O-R^{10}$, $-S-R^{10}$, $-S(O)-R^{10}$, $-SO_2-R^{10}$, $-SO_2-OR^{10}$, $-SO_2-NR^{11}R^{10}$, $-CO-OR^{10}$, $-CO-NR^{11}R^{10}$, $-O-SO_2-R^{10}$, $-N(R^{11})-SO_2-R^{10}$, $-NR^{11}R^{10}$, $-NR^{11}R^{10}$, $-NH-P(O)(R^{11})(OR^{10})$,

R⁵ represents nitro, cyano, fluorine, chlorine or bromine or represents straight-chain or branched halogenoalkyl having 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms of fluorine, chlorine or bromine, and

X represents oxygen or sulphur, where

 R^8 represents hydrogen, straight-chain or branched alkyl having from 1 to 6 carbon atoms or straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms of fluorine, chlorine or bromine, and furthermore represents a radical of the formula $-CO-R^{12}$ or a radical of the formula $-S(O)_n-R^{12}$,

 R^9 represents straight-chain or branched alkyl having from 1 to 6 carbon atoms or straight-chain or branched halogeno-alkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms of fluorine, chlorine or bromine, and furthermore represents a radical of the formula $-CO-R^{12}$ or a radical of the formula $-S(O)_n-R^{12}$,

R¹⁰ represents hydrogen,

R¹⁰ furthermore represents straight-chain or branched alkyl having from 1 to 12 carbon atoms which is optionally substituted once or twice by identical or different substituents of cyano, carboxyl, carbamoyl, in each case straight-chain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, N-alkylaminocarbonyl, N,N-dialkylamino-

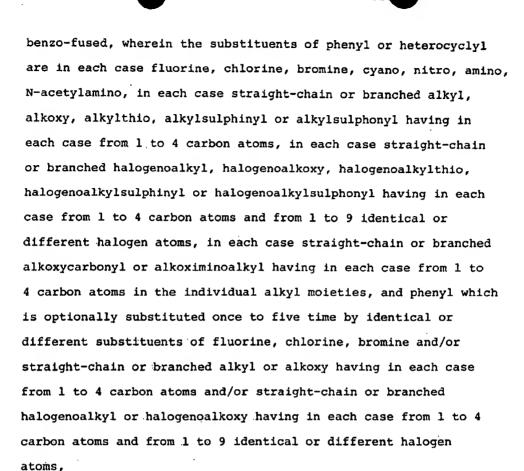
carbonyl or alkylsulphonylaminocarbonyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, or heterocyclyl, the heterocyclyl radical being a five- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur,

R¹⁰ furthermore represents straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms of fluorine, chlorine and/or bromine.

R¹⁰ furthermore represents alkenyl or alkinyl having in each case from 2 to 6 carbon atoms, which are in each case optionally substituted once to three times by identical or different halogen atoms of fluorine, chlorine and/or bromine,

R¹⁰ furthermore represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once to three times by identical or different substituents of fluorine, chlorine, bromine and/or straight-chain or branched alkyl having from 1 to 3 carbon atoms,

R¹⁰ furthermore represents phenylalkyl or phenyl having optionally from 1 to 3 carbon atoms in the straight-chain or branched akyl moiety, which are in each case optionally substituted in the phenyl moiety once to five times by identical or different substituents, or represents a saturated or unsaturated, five- to seven-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur, which is optionally substituted once to three times by identical or different substituents and/or is



R¹¹ represents hydrogen,

R¹¹ furthermore represents straight-chain or branched alkyl having from 1 to 12 carbon atoms which is optionally substituted once or twice by identical or different substituents of cyano, carboxyl, carbamoyl, in each case straight-chain or branched alkoxy, alkoxyalkoxy, alkylthio, alkylsulphinyl, alkylsulphonyl, alkoxycarbonyl, N-alkylaminocarbonyl, N,N-dialkylaminocarbonyl or alkylsulphonylaminocarbonyl having in each case from 1 to 6 carbon atoms in the individual alkyl moieties, or

heterocyclyl, the heterocyclyl radical being a five- to sevenmembered, optionally benzo-fused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur,

R¹¹ furthermore represents straight-chain or branched halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms of fluorine, chlorine and/or bromine,

R¹¹ furthermore represents alkenyl or alkinyl having in each case from 2 to 6 carbon atoms, which are in each case optionally substituted once to three times by identical or different halogen atoms of fluorine, chlorine and/or bromine,

R¹¹ furthermore represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once to three times by identical or different substituents of fluorine, chlorine, bromine and/or straight-chain or branched alkyl having from 1 to 3 carbon atoms,

R¹¹ furthermore represents phenylalkyl or phenyl having optionally from 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the phenyl moiety once to five times by identical or different substituents, wherein the substituents of phenyl are in each case fluorine, chlorine, bromine, cyano, nitro, amino, N-acetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 4 carbon atoms, in each case straight-chain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in

each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 4 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once to five times by identical or different substituents of fluorine, chlorine, bromine and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 4 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms;

R¹² represents straight-chain or branched alkyl having from 1 to 12 carbon atoms which is optionally substituted once or twice by identical or different substituents of cycloalkyl having from 3 to 7 carbon atoms or heterocyclyl, the heterocyclyl radical being a five- to seven-membered, optionally benzo-fused, saturated or unsaturated heterocycle having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur,

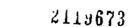
R¹² furthermore represents halogenoalkyl having from 1 to 6 carbon atoms and from 1 to 13 identical or different halogen atoms of fluorine, chlorine and/or bromine,

R¹² furthermore represents cycloalkyl having from 3 to 7 carbon atoms which is optionally substituted once to three times by identical or different substituents of fluorine, chlorine, bromine and/or straight-chain or branched alkyl having from 1 to 3 carbon atoms,

 R^{12} furthermore represents phenylalkyl or phenyl having

optionally from 1 to 3 carbon atoms in the straight-chain or branched alkyl moiety, which are in each case optionally substituted in the phenyl moiety once to five times by identical or different substituents, or represents a saturated or unsaturated, five- to seven-membered heterocyclyl radical having from 1 to 3 identical or different hetero atoms of nitrogen, oxygen and/or sulphur, which is optionally substituted once to three times by identical or different substituents and/or is benzo-fused, wherein the substituents of phenyl or heterocyclyl are in each case fluorine, chlorine, bromine, cyano, nitro, amino, N-acetylamino, in each case straight-chain or branched alkyl, alkoxy, alkylthio, alkylsulphinyl or alkylsulphonyl having in each case from 1 to 4 carbon atoms, in each case straightchain or branched halogenoalkyl, halogenoalkoxy, halogenoalkylthio, halogenoalkylsulphinyl or halogenoalkylsulphonyl having in each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms, in each case straight-chain or branched alkoxycarbonyl or alkoximinoalkyl having in each case from 1 to 4 carbon atoms in the individual alkyl moieties, and phenyl which is optionally substituted once to five times by identical or different substituents of fluorine, chlorine, bromine and/or straight-chain or branched alkyl or alkoxy having in each case from 1 to 4 carbon atoms and/or straight-chain or branched halogenoalkyl or halogenoalkoxy having in each case from 1 to 4 carbon atoms and from 1 to 9 identical or different halogen atoms and

n represents a number 0, 1 or 2.





4. The compound 1-(4-cyano-2,5-difluoropheny1)-3-methy1-4-dimethylamino-1,2,4-triazolin-5-one of the formula

5. The compound 1-(4-cyano-2,5-difluoropheny1)-3-methy1-4-(N-methylamino)-1,2,4-triazolin-5-one of the formula

6. A compound according to claim 1 wherein ${\tt R}^1 \text{ is methyl, } {\tt R}^2 \text{ is } -{\tt N(CH}_3)_2, \ {\tt R}^3 \text{ is fluorine, } {\tt R}^4 \text{ is hydrogen, } {\tt R}^5 \text{ is cyano, X is oxygen and } {\tt R}^7 \text{ is } {\tt CH}_3 \\ {\tt o-ch-c-c-ch}.$

- 7. A compound according to claim 1 wherein R^1 is methyl, R^2 is $-N(CH_3)_2$, R^3 is fluorine, R^4 is hydrogen, R^5 is cyano, X is oxygen and R^7 is NH-SO₂-CH₃.
- 8. A herbicidal composition comprising a herbicidally effective amount of a compound according to any one of claims 1 to 7 in admixture with a suitable carrier or diluent.
- 9. A herbicidal composition comprising a herbicidally effective amount of a compound according to any one of claims 1 to 7 in admixture with a solid diluent or carrier, a liquified normally gaseous diluent or carrier, or a liquid diluent or carrier containing a surface active agent.
- 10. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a compound according to any one of claims 1 to 7.
- 11. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a composition containing a compound according to any one of claims 1 to 7 in admixture with a suitable carrier or diluent.
- 12. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a composition containing between 0.1 and 95% by weight of a compound according to any one of claims 1 to 7 in admixture with a suitable carrier or diluent.



- 13. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a composition containing between 0.5 and 90% by weight of a compound according to any one of claims 1 to 7 in admixture with a suitable carrier or diluent.
- 14. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a compound according to any one of claims 1 to 7 wherein the compound is applied as a pre-emergence herbicide.
- 15. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a compound according to any one of claims 1 to 7 wherein the compound is applied as a post-emergence herbicide.
- 16. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a compound according to any one of claims 1 to 7 wherein the compound is applied to an area of cultivation at a rate of between 0.01 and 10 kg/ha.
- 17. A method of combating weeds which comprises applying to the weeds, or to a habitat thereof, a herbicidally effective amount of a compound according to any one of claims 1 to 7 wherein the compound is applied to an area of cultivation at a rate of between 0.05 and 5 kg/ha.
- 18. A process for preparing a compound of formula (I) as defined in claim 1 and R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 and X are as



defined in claim 1, which process comprises:

a) reacting a lH-triazolinone of the formula (II)

in which

 ${\mbox{R}}^1, \ {\mbox{R}}^2$ and X have the meaning given above, with a halogenobenzene derivative of the formula (III)

$$\begin{array}{c}
R^3 \\
R^4 \\
R^5
\end{array}$$
(III)

in which

 ${\bf R}^3$, ${\bf R}^4$, ${\bf R}^5$, ${\bf R}^6$ and ${\bf R}^7$ have the meanings given above and ${\bf Hal}^1$ represents halogen, or

b) reacting a substituted 1-aryltriazolinone of the formula (Ia)

$$R^{1}$$
 N
 N
 X
 R^{7}
 R^{3}
 R^{6}
 R^{5}
 R^{1}
 R^{2}
 R^{3}
 R^{3}
 R^{6}
 R^{5}

- 120 -

in which

 ${\rm R}^1,~{\rm R}^2,~{\rm R}^3,~{\rm R}^6,~{\rm R}^5,~{\rm R}^7$ and X have the meanings given above and

 ${\rm Hal}^2$ represents halogen, with a nucleophile of the formula (IV)

$$H-R^{13}$$
 (IV)

in which

 $\rm R^{13}$ represents a radical of the formula -0-R 10 , -S-R 10 or -NR $^{11}\rm R^{10}$, where R 10 and R 11 have the meanings given above, or

c) reacting a substituted triazolinone of the formula

(V)

in which

 R^1 , R^3 , R^4 , R^5 , R^6 , R^7 and X have the meanings given above, with an alkylating, acylating or sulphonylating agent of the formula (VI)

$$R^9-E$$
 (VI)

in which

 ${\ensuremath{\mathtt{R}}}^9$ has the meaning given above and ${\ensuremath{\mathtt{E}}}$ represents an electron-attracting leaving group, or

d) reacting a 4-alkylideneimino-triazolinone of the formula (VII)

$$\begin{array}{c}
R^{1} \\
N \\
N \\
N
\end{array}$$

$$\begin{array}{c}
R^{14} \\
R^{15} \\
R^{3} \\
R^{6}
\end{array}$$
(VII)

in which

 ${\bf R}^1,~{\bf R}^3,~{\bf R}^4,~{\bf R}^5,~{\bf R}^6,~{\bf R}^7$ and X have the meanings given above,

 $\ensuremath{\text{R}^{14}}$ represents hydrogen or alkyl and $\ensuremath{\text{R}^{15}}$ represents alkyl or alkoxy.

- 19. A process for preparing a herbicidal composition which comprises admixing a compound of formula (I) as defined in any one of claims 1 to 7 together with an extender or surface active agent.
- 20. A substituted 1-aryltriazolinone of the formula (Ia)



in which

 R^1 , R^2 , R^3 , R^5 , R^6 , R^7 and X are as defined in claim 1

and

Hal² represents halogen.

21. A substituted triazolinone of the formula (V)

in which

 R^{1} , R^{3} , R^{4} , R^{5} , R^{6} , R^{7} and X are as defined in claim 1.

FETHERSTONHAUGH & CO. OTTAWA, CANADA

PATENT AGENTS